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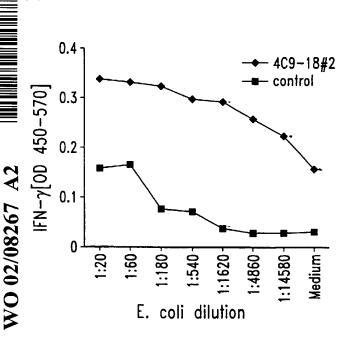
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(54) Title: COMPOUNDS AND METHODS FOR TREATMENT AND DIAGNOSIS OF CHLAMYDIAL INFECTION



(57) Abstract: Compounds and methods for the diagnosis and treatment of Chlamydial infection are disclosed. The compounds provided include polypeptides that contain at least one antigenic portion of a *Chlamydia* antigen and DNA sequences encoding such polypeptides. Pharmaceutical compositions and vaccines comprising such polypeptides or DNA sequences are also provided, together with antibodies directed against such polypeptides. Diagnostic kits containing such polypeptides or DNA sequences and a suitable detection reagent may be used for the detection of Chlamydial infection in patients and in biological samples.

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COMPOUNDS AND METHODS FOR TREATMENT AND DIAGNOSIS OF CHLAMYDIAL INFECTION

TECHNICAL FIELD

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The present invention relates generally to the detection and treatment of Chlamydial infection. In particular, the invention is related to polypeptides comprising a *Chlamydia* antigen and the use of such polypeptides for the serodiagnosis and treatment of Chlamydial infection.

10 BACKGROUND OF THE INVENTION

Chlamydiae are intracellular bacterial pathogens that are responsible for a wide variety of important human and animal infections. Chlamydia trachomatis is one of the most common causes of sexually transmitted diseases and can lead to pelvic inflammatory disease (PID), resulting in tubal obstruction and infertility. Chlamydia trachomatis may also play a role in male infertility. In 1990, the cost of treating PID in the US was estimated to be \$4 billion. Trachoma, due to ocular infection with Chlamydia trachomatis, is the leading cause of preventable blindness worldwide. Chlamydia pneumonia is a major cause of acute respiratory tract infections in humans and is also believed to play a role in the pathogenesis of atherosclerosis and, in particular, coronary heart disease. Individuals with a high titer of antibodies to Chlamydia pneumonia have been shown to be at least twice as likely to suffer from coronary heart disease as seronegative individuals. Chlamydial infections thus constitute a significant health problem both in the US and worldwide.

Chlamydial infection is often asymptomatic. For example, by the time a woman seeks medical attention for PID, irreversible damage may have already occurred resulting in infertility. There thus remains a need in the art for improved vaccines and pharmaceutical compositions for the prevention and treatment of *Chlamydia* infections. The present invention fulfills this need and further provides other related advantages.

30 SUMMARY OF THE INVENTION

The present invention provides compositions and methods for the diagnosis and therapy of *Chlamydia* infection. In one aspect, the present invention

2

provides polypeptides comprising an immunogenic portion of a *Chlamydia* antigen, or a variant of such an antigen. Certain portions and other variants are immunogenic, such that the ability of the variant to react with antigen-specific antisera is not substantially diminished. Within certain embodiments,, the polypeptide comprises an amino acid sequence encoded by a polynucleotide sequence selected from the group consisting of (a) a sequence of SEQ ID NO: 358-361, 366-385, 406-430, 455-489, 516-517, 523-559, and 582-596; (b) the complements of said sequences; and (c) sequences that hybridize to a sequence of (a) or (b) under moderate to highly stringent conditions. In specific embodiments, the polypeptides of the present invention comprise at least a portion of a *Chlamydial* protein that includes an amino acid sequence selected from the group consisting of sequences recited in SEQ ID NO:362-365, 386-405, 431-454, 490-515, 518-522, 560-581, and 597-599 and variants thereof.

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The present invention further provides polynucleotides that encode a polypeptide as described above, or a portion thereof (such as a portion encoding at least 15 amino acid residues of a *Chlamydial* protein), expression vectors comprising such polynucleotides and host cells transformed or transfected with such expression vectors.

In a related aspect, polynucleotide sequences encoding the above polypeptides, recombinant expression vectors comprising one or more of these polynucleotide sequences and host cells transformed or transfected with such expression vectors are also provided.

In another aspect, the present invention provides fusion proteins comprising an inventive polypeptide, or, alternatively, an inventive polypeptide and a known *Chlamydia* antigen, as well as polynucleotides encoding such fusion proteins, in combination with a physiologically acceptable carrier or immunostimulant for use as pharmaceutical compositions and vaccines thereof.

The present invention further provides pharmaceutical compositions that comprise: (a) an antibody, both polyclonal and monoclonal, or antigen-binding fragment thereof that specifically binds to a *Chlamydial* protein; and (b) a physiologically acceptable carrier. Within other aspects, the present invention provides pharmaceutical compositions that comprise one or more *Chlamydia* polypeptides disclosed herein, e.g., a polypeptide according to SEQ ID NO:362-365, 386-405, 431-454, 490-515, 518-522, 560-581, and 597-599, or a polynucleotide molecule encoding

3

such a polypeptide, such as a polynucleotide according to SEQ ID NO:358-361, 366-385, 406-430, 455-489, 516-517, 523-559, and 582-596, and a physiologically acceptable carrier. The invention also provides vaccines for prophylactic and therapeutic purposes comprising one or more of the disclosed polypeptides and an immunostimulant, as defined herein, together with vaccines comprising one or more polynucleotide sequences encoding such polypeptides and an immunostimulant.

In yet another aspect, methods are provided for inducing protective immunity in a patient, comprising administering to a patient an effective amount of one or more of the above pharmaceutical compositions or vaccines.

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In yet a further aspect, methods for the treatment of Chlamydia infection in a patient are provided, the methods comprising obtaining peripheral blood mononuclear cells (PBMC) from the patient, incubating the PBMC with a polypeptide of the present invention (or a polynucleotide that encodes such a polypeptide) to provide incubated T cells and administering the incubated T cells to the patient. The present invention additionally provides methods for the treatment of Chlamydia infection that comprise incubating antigen presenting cells with a polypeptide of the present invention (or a polynucleotide that encodes such a polypeptide) to provide incubated antigen presenting cells and administering the incubated antigen presenting cells to the patient. Proliferated cells may, but need not, be cloned prior to administration to the patient. In certain embodiments, the antigen presenting cells are selected from the group consisting of dendritic cells, macrophages, monocytes, B-cells, and fibroblasts. Compositions for the treatment of Chlamydia infection comprising T cells or antigen presenting cells that have been incubated with a polypeptide or polynucleotide of the present invention are also provided. Within related aspects, vaccines are provided that comprise: (a) an antigen presenting cell that expresses a polypeptide as described above and (b) an immunostimulant.

The present invention further provides, within other aspects, methods for removing *Chlamydial*-infected cells from a biological sample, comprising contacting a biological sample with T cells that specifically react with a *Chlamydial* protein, wherein the step of contacting is performed under conditions and for a time sufficient to permit the removal of cells expressing the protein from the sample.

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Within related aspects, methods are provided for inhibiting the development of *Chlamydial* infection in a patient, comprising administering to a patient a biological sample treated as described above. In further aspects of the subject invention, methods and diagnostic kits are provided for detecting *Chlamydia* infection in a patient. In one embodiment, the method comprises: (a) contacting a biological sample with at least one of the polypeptides or fusion proteins disclosed herein; and (b) detecting in the sample the presence of binding agents that bind to the polypeptide or fusion protein, thereby detecting *Chlamydia* infection in the biological sample. Suitable biological samples include whole blood, sputum, serum, plasma, saliva, cerebrospinal fluid and urine. In one embodiment, the diagnostic kits comprise one or more of the polypeptides or fusion proteins disclosed herein in combination with a detection reagent. In yet another embodiment, the diagnostic kits comprise either a monoclonal antibody or a polyclonal antibody that binds with a polypeptide of the present invention.

The present invention also provides methods for detecting *Chlamydia* infection comprising: (a) obtaining a biological sample from a patient; (b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, at least one of the oligonucleotide primers being specific for a polynucleotide sequence disclosed herein; and (c) detecting in the sample a polynucleotide sequence that amplifies in the presence of the oligonucleotide primers. In one embodiment, the oligonucleotide primer comprises at least about 10 contiguous nucleotides of a polynucleotide sequence peptide disclosed herein, or of a sequence that hybridizes thereto.

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In a further aspect, the present invention provides a method for detecting *Chlamydia* infection in a patient comprising: (a) obtaining a biological sample from the patient; (b) contacting the sample with an oligonucleotide probe specific for a polynucleotide sequence disclosed herein; and (c) detecting in the sample a polynucleotide sequence that hybridizes to the oligonucleotide probe. In one embodiment, the oligonucleotide probe comprises at least about 15 contiguous nucleotides of a polynucleotide sequence disclosed herein, or a sequence that hybridizes thereto.

These and other aspects of the present invention will become apparent upon reference to the following detailed description. All references disclosed herein are

hereby incorporated by reference in their entirety as if each was incorporated individually.

SEQUENCE IDENTIFIERS

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5 SEQ ID NO: 1 is the determined DNA sequence for the *C. trachomatis* clone 1-B1-66.

SEQ ID NO: 2 is the determined DNA sequence for the *C. trachomatis* clone 4-D7-28.

SEQ ID NO: 3 is the determined DNA sequence for the *C. trachomatis* clone 3-G3-10.

SEQ ID NO: 4 is the determined DNA sequence for the *C. trachomatis* clone 10-C10-31.

SEQ ID NO: 5 is the predicted amino acid sequence for 1-B1-66.

SEQ ID NO: 6 is the predicted amino acid sequence for 4-D7-28.

SEQ ID NO: 7 is a first predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 8 is a second predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 9 is a third predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 10 is a fourth predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 11 is a fifth predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 12 is the predicted amino acid sequence for 10-C10-31.

SEQ ID NO: 13 is the amino acid sequence of the synthetic peptide 1-B1-66/48-67.

SEQ ID NO: 14 is the amino acid sequence of the synthetic peptide 1-B1-66/58-77.

SEQ ID NO: 15 is the determined DNA sequence for the *C. trachomatis* serovar LGV II clone 2C7-8

SEQ ID NO: 16 is a DNA sequence of a putative open reading frame from a region of the C. trachomatis serovar D genome to which 2C7-8 maps

SEQ ID NO: 17 is the predicted amino acid sequence encoded by the 30 DNA sequence of SEQ ID NO: 16

SEQ ID NO: 18 is the amino acid sequence of the synthetic peptide CtC7.8-12

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SEQ ID NO: 19 is the amino acid sequence of the synthetic peptide CtC7.8-13

SEQ ID NO: 20 is the predicted amino acid sequence encoded by a second putative open reading from C. trachomatis serovar D

SEQ ID NO: 21 is the determined DNA sequence for clone 4C9-18 from C. trachomatis LGV II

SEQ ID NO: 22 is the determined DNA sequence homologous to Lipoamide Dehydrogenase from *C. trachomatis* LGV II

SEQ ID NO: 23 is the determined DNA sequence homologous to Hypothetical protein from C. trachomatis LGV II

SEQ ID NO: 24 is the determined DNA sequence homologous to Ubiquinone Mehtyltransferase from C. trachomatis LGV II

SEQ ID NO: 25 is the determined DNA sequence for clone 4C9-18#2 BL21 pLysS from C. trachomatis LGV II

SEQ ID NO: 26 is the predicted amino acid sequence for 4C9-18#2 from C. trachomatis LGV II

SEQ ID NO: 27 is the determined DNA sequence for Cp-SWIB from C. pneumonia strain TWAR

SEQ ID NO: 28 is the predicted amino acid sequence for Cp-SWIB from 20 *C. pneumonia* strain TWAR

SEQ ID NO: 29 is the determined DNA sequence for Cp-S13 (CT509) from *C. pneumonia* strain TWAR

SEQ ID NO: 30 is the predicted amino acid sequence for Cp-S13 from C. pneumonia strain TWAR

SEQ ID NO: 31 is the amino acid sequence for a 10mer consensus peptide from CtC7.8-12 and CtC7.8-13

SEQ ID NO: 32 is the predicted amino acid sequence for clone 2C7-8 from $\it C. trachomatis LGV II$

SEQ ID NO: 33 is the DNA sequence corresponding to nucleotides 597304-597145 of the *C. trachomatis* serovar D genome (NCBI, BLASTN search), which shows homology to clone 2C7-8

SEQ ID NO: 34 is the predicted amino acid sequence encoded by the sequence of SEQ ID NO: 33

SEQ ID NO: 35 is the DNA sequence for C.p. SWIB Nde (5' primer) from C. pneumonia

SEQ ID NO: 36 is the DNA sequence for C.p. SWIB EcoRI (3' primer) from C. pneumonia

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SEQ ID NO: 37 is the DNA sequence for C.p. S13 Nde (5' primer) from C. pneumonia

SEQ ID NO: 38 is the DNA sequence for C.p. S13 EcoRI (3' primer) from C. pneumonia

SEQ ID NO: 39 is the amino acid sequence for CtSwib 52-67 peptide from C. trachomatis LGV II

SEQ ID NO: 40 is the amino acid sequence for CpSwib 53-68 peptide from C. pneumonia

SEQ ID NO: 41 is the amino acid sequence for HuSwib 288-302 peptide from Human SWI domain

SEQ ID NO: 42 is the amino acid sequence for CtSWI-T 822-837 peptide from the topoisomerase-SWIB fusion of *C. trachomatis*

SEQ ID NO: 43 is the amino acid sequence for CpSWI-T 828-842 peptide from the topoisomerase-SWIB fusion of *C. pneumonia*

SEQ ID NO: 44 is a first determined DNA sequence for the C. trachomatis LGV II clone 19783.3, jen.seq(1>509)CTL2#11-3', representing the 3' end.

SEQ ID NO: 45 is a second determined DNA sequence for the C. trachomatis LGV II clone 19783.4,jen.seq(1>481)CTL2#11-5', representing the 5' end.

SEQ ID NO: 46 is the determined DNA sequence for the *C. trachomatis* LGV II clone19784CTL2_12consensus.seq(1>427)CTL2#12.

SEQ ID NO: 47 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19785.4,jen.seq(1>600)CTL2#16-5', representing the 5' end.

SEQ ID NO: 48 is a first determined DNA sequence for the C. trachomatis LGV II clone 19786.3, jen.seq(1>600)CTL2#18-3', representing the 3' end.

SEQ ID NO: 49 is a second determined DNA sequence for the C. trachomatis LGV II clone 19786.4, jen. seq(1>600) CTL2#18-5', representing the 5' end.

SEQ ID NO: 50 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19788CTL2_21consensus.seq(1>406)CTL2#21.

SEQ ID NO: 51 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19790CTL2_23consensus.seq(1>602)CTL2#23.

5 SEQ ID NO: 52 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19791CTL2_24consensus.seq(1>145)CTL2#24.

SEQ ID NO: 53 is the determined DNA sequence for the *C. trachomatis* LGV II clone CTL2#4.

SEQ ID NO: 54 is the determined DNA sequence for the *C. trachomatis* 10 LGV II clone CTL2#8b.

SEQ ID NO: 55 is the determined DNA sequence for the *C. trachomatis* LGV II clone15-G1-89, sharing homology to the lipoamide dehydrogenase gene CT557.

SEQ ID NO: 56 is the determined DNA sequence for the *C. trachomatis* LGV II clone 14-H1-4, sharing homology to the thiol specific antioxidant gene CT603.

SEQ ID NO: 57 is the determined DNA sequence for the *C. trachomatis* LGV II clone 12-G3-83, sharing homology to the hypothetical protein CT622.

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SEQ ID NO: 58 is the determined DNA sequence for the *C. trachomatis* LGV II clone 12-B3-95, sharing homology to the lipoamide dehydrogenase gene CT557.

SEQ ID NO: 59 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-H4-28, sharing homology to the dnaK gene CT396.

SEQ ID NO: 60 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-H3-68, sharing partial homology to the PGP6-D virulence protein and L1 ribosomal gene CT318.

SEQ ID NO: 61 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-G1-34, sharing partial homology to the malate dehydrogenase gene CT376 and to the glycogen hydrolase gene CT042.

SEQ ID NO: 62 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-G10-46, sharing homology to the hypothetical protein CT610.

SEQ ID NO: 63 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-C12-91, sharing homology to the OMP2 gene CT443.

SEQ ID NO: 64 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-A3-93, sharing homology to the HAD superfamily gene CT103.

SEQ ID NO: 65 is the determined amino acid sequence for the *C. trachomatis* LGV II clone 14-H1-4, sharing homology to the thiol specific antioxidant gene CT603.

SEQ ID NO: 66 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#9.

SEQ ID NO: 67 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#7.

SEQ ID NO: 68 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#6.

SEQ ID NO: 69 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#5.

SEQ ID NO: 70 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#2.

SEQ ID NO: 71 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#1.

SEQ ID NO: 72 is a first determined DNA sequence for the *C. trachomatis* LGV II clone 23509.2CtL2#3-5', representing the 5' end.

SEQ ID NO: 73 is a second determined DNA sequence for the C. trachomatis LGV II clone 23509.1CtL2#3-3', representing the 3' end.

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SEQ ID NO: 74 is a first determined DNA sequence for the *C. trachomatis* LGV II clone 22121.2CtL2#10-5', representing the 5' end.

SEQ ID NO: 75 is a second determined DNA sequence for the C. trachomatis LGV II clone 22121.1CtL2#10-3', representing the 3' end.

SEQ ID NO: 76 is the determined DNA sequence for the *C. trachomatis*. LGV II clone 19787.6CtL2#19-5', representing the 5' end.

SEQ ID NO: 77 is the determined DNA sequence for the *C. pneumoniae* LGV II clone CpS13-His.

SEQ ID NO: 78 is the determined DNA sequence for the *C. pneumoniae* LGV II clone Cp SWIB-His.

SEQ ID NO: 79 is the determined DNA sequence for the *C. trachomatis* LGV II clone 23-G7-68, sharing partial homology to the L11, L10 and L1 ribosomal protein.

SEQ ID NO: 80 is the determined DNA sequence for the *C. trachomatis* LGV II clone 22-F8-91, sharing homology to the pmpC gene.

SEQ ID NO: 81 is the determined DNA sequence for the *C. trachomatis* LGV II clone 21-E8-95, sharing homology to the CT610-CT613 genes.

SEQ ID NO: 82 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19-F12-57, sharing homology to the CT858 and recA genes.

SEQ ID NO: 83 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19-F12-53, sharing homology to the CT445 gene encoding glutamyl tRNA synthetase.

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SEQ ID NO: 84 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19-A5-54, sharing homology to the cryptic plasmid gene.

SEQ ID NO: 85 is the determined DNA sequence for the *C. trachomatis* LGV II clone 17-E11-72, sharing partial homology to the OppC_2 and pmpD genes.

SEQ ID NO: 86 is the determined DNA sequence for the *C. trachomatis* LGV II clone 17-C1-77, sharing partial homology to the CT857 and CT858 open reading frames.

SEQ ID NO: 87 is the determined DNA sequence for the *C. trachomatis* LGV II clone 15-H2-76, sharing partial homology to the pmpD and SycE genes, and to the CT089 ORF.

SEQ ID NO: 88 is the determined DNA sequence for the *C. trachomatis* LGV II clone 15-A3-26, sharing homology to the CT858 ORF.

SEQ ID NO: 89 is the determined amino acid sequence for the C. pnuemoniae clone Cp_SWIB-His.

SEQ ID NO: 90 is the determined amino acid sequence for the C. trachomatis LGV II clone CtL2 LPDA FL.

SEQ ID NO: 91 is the determined amino acid sequence for the C. pnuemoniae clone CpS13-His.

SEQ ID NO: 92 is the determined amino acid sequence for the C. trachomatis LGV II clone CtL2 TSA FL.

SEQ ID NO: 93 is the amino acid sequence for Ct-Swib 43-61 peptide from C. trachomatis LGV II.

SEQ ID NO: 94 is the amino acid sequence for Ct-Swib 48-67 peptide from C. trachomatis LGV II.

5 SEQ ID NO: 95 is the amino acid sequence for Ct-Swib 52-71 peptide from *C. trachomatis* LGV II.

SEQ ID NO: 96 is the amino acid sequence for Ct-Swib 58-77 peptide from C. trachomatis LGV II.

SEQ ID NO: 97 is the amino acid sequence for Ct-Swib 63-82 peptide 10 from C. trachomatis LGV II.

SEQ ID NO: 98 is the amino acid sequence for Ct-Swib 51-66 peptide from C. trachomatis LGV II.

SEQ ID NO: 99 is the amino acid sequence for Cp-Swib 52-67 peptide from *C. pneumonia*.

SEQ ID NO: 100 is the amino acid sequence for Cp-Swib 37-51 peptide from *C. pneumonia*.

SEQ ID NO: 101 is the amino acid sequence for Cp-Swib 32-51 peptide from *C. pneumonia*.

SEQ ID NO: 102 is the amino acid sequence for Cp-Swib 37-56 peptide 20 from *C. pneumonia*.

SEQ ID NO: 103 is the amino acid sequence for Ct-Swib 36-50 peptide from C. trachomatis.

SEQ ID NO: 104 is the amino acid sequence for Ct-S13 46-65 peptide from C. trachomatis.

SEQ ID NO: 105 is the amino acid sequence for Ct-S13 60-80 peptide from C. trachomatis.

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SEQ ID NO: 106 is the amino acid sequence for Ct-S13 1-20 peptide from C. trachomatis.

SEQ ID NO: 107 is the amino acid sequence for Ct-S13 46-65 peptide from C. trachomatis.

SEQ ID NO: 108 is the amino acid sequence for Ct-S13 56-75 peptide from C. trachomatis.

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SEQ ID NO: 109 is the amino acid sequence for Cp-S13 56-75 peptide from *C. pneumoniae*.

SEQ ID NO: 110 is the determined DNA sequence for the *C. trachomatis* LGV II clone 21-G12-60, containing partial open reading frames for hypothetical proteins CT875, CT229 and CT228.

SEQ ID NO: 111 is the determined DNA sequence for the C. trachomatis LGV II clone 22-B3-53, sharing homology to the CT110 ORF of GroEL.

SEQ ID NO: 112 is the determined DNA sequence for the *C. trachomatis* LGV II clone 22-A1-49, sharing partial homology to the CT660 and CT659 ORFs.

SEQ ID NO: 113 is the determined DNA sequence for the *C. trachomatis* LGV II clone 17-E2-9, sharing partial homology to the CT611 and CT 610 ORFs.

SEQ ID NO: 114 is the determined DNA sequence for the C. trachomatis LGV II clone 17-C10-31, sharing partial homology to the CT858 ORF.

SEQ ID NO: 115 is the determined DNA sequence for the *C. trachomatis* LGV II clone 21-C7-8, sharing homology to the dnaK-like gene.

SEQ ID NO: 116 is the determined DNA sequence for the *C. trachomatis* LGV II clone 20-G3-45, containing part of the pmpB gene CT413.

SEQ ID NO: 117 is the determined DNA sequence for the *C. trachomatis* LGV II clone 18-C5-2, sharing homology to the S1 ribosomal protein ORF.

SEQ ID NO: 118 is the determined DNA sequence for the C. trachomatis LGV II clone 17-C5-19, containing part of the ORFs for CT431 and CT430.

SEQ ID NO: 119 is the determined DNA sequence for the *C. trachomatis* LGV II clone 16-D4-22, contains partial sequences of ORF3 and ORF4 of the plasmid for growth within mammalian cells.

SEQ ID NO: 120 is the determined full-length DNA sequence for the *C. trachomatis* serovar LGV II Cap1 gene CT529.

SEQ ID NO: 121 is the predicted full-length amino acid sequence for the *C. trachomatis* serovar LGV II Cap1 gene CT529.

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SEQ ID NO: 122 is the determined full-length DNA sequence for the *C. trachomatis* serovar E Cap1 gene CT529.

SEQ ID NO: 123 is the predicted full-length amino acid sequence for the C. trachomatis serovar E Cap1 gene CT529.

SEQ ID NO: 124 is the determined full-length DNA sequence for the C. trachomatis serovar 1A Cap1 gene CT529.

SEQ ID NO: 125 is the predicted full-length amino acid sequence for the C. trachomatis serovar 1A Cap1 gene CT529.

SEQ ID NO: 126 is the determined full-length DNA sequence for the C.

10 trachomatis serovar G Cap1 gene CT529.

SEQ ID NO: 127 is the predicted full-length amino acid sequence for the *C. trachomatis* serovar G Cap1 gene CT529.

SEQ ID NO: 128 is the determined full-length DNA sequence for the *C. trachomatis* serovar F1 NII Cap1 gene CT529.

SEQ ID NO: 129 is the predicted full-length amino acid sequence for the C. trachomatis serovar F1 NII Cap1 gene CT529.

SEQ ID NO: 130 is the determined full-length DNA sequence for the *C. trachomatis* serovar L1 Cap1 gene CT529.

SEQ ID NO: 131 is the predicted full-length amino acid sequence for the 20 C. trachomatis serovar L1 Cap1 gene CT529.

SEQ ID NO: 132 is the determined full-length DNA sequence for the *C. trachomatis* serovar L3 Cap1 gene CT529.

SEQ ID NO: 133 is the predicted full-length amino acid sequence for the C. trachomatis serovar L3 Cap1 gene CT529.

SEQ ID NO: 134 is the determined full-length DNA sequence for the C. trachomatis serovar Ba Cap1 gene CT529.

SEQ ID NO: 135 is the predicted full-length amino acid sequence for the *C. trachomatis* serovar Ba Cap1 gene CT529.

SEQ ID NO: 136 is the determined full-length DNA sequence for the C.

trachomatis serovar MOPN Cap1 gene CT529.

SEQ ID NO: 137 is the predicted full-length amino acid sequence for the C. trachomatis serovar MOPN Cap1 gene CT529.

SEQ ID NO: 138 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #124-139 of *C. trachomatis* serovar L2.

SEQ ID NO: 139 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #132-147 of *C. trachomatis* serovar L2.

SEQ ID NO: 140 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #138-155 of *C. trachomatis* serovar L2.

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SEQ ID NO: 141 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #146-163 of *C. trachomatis* serovar L2.

SEQ ID NO: 142 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #154-171 of *C. trachomatis* serovar L2.

SEQ ID NO: 143 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #162-178 of *C. trachomatis* serovar L2.

SEQ ID NO: 144 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #138-147 of *C. trachomatis* serovar L2.

SEQ ID NO: 145 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #139-147 of *C. trachomatis* serovar L2.

SEQ ID NO: 146 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #140-147 of *C. trachomatis* serovar L2.

SEQ ID NO: 147 is the determined amino acid sequence for the Cap1

20 CT529 ORF peptide #138-146 of *C. trachomatis* serovar L2.

SEQ ID NO: 148 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #138-145 of *C. trachomatis* serovar L2.

SEQ ID NO: 149 is the determined amino acid sequence for the Cap1 CT529 ORF peptide # F140->I of *C. trachomatis* serovar L2.

SEQ ID NO: 150 is the determined amino acid sequence for the Cap1 CT529 ORF peptide ##S139>Ga of *C. trachomatis* serovar L2.

SEQ ID NO: 151 is the determined amino acid sequence for the Cap1 CT529 ORF peptide ##S139>Gb of *C. trachomatis* serovar L2.

SEQ ID NO: 152 is the determined amino acid sequence for the peptide # 2 C7.8-6 of the 216aa ORF of C. trachomatis serovar L2.

SEQ ID NO: 153 is the determined amino acid sequence for the peptide # 2 C7.8-7 of the 216aa ORF of *C. trachomatis* serovar L2.

SEQ ID NO: 154 is the determined amino acid sequence for the peptide #2 C7.8-8 of the 216aa ORF of *C. trachomatis* servoar L2.

SEQ ID NO: 155 is the determined amino acid sequence for the peptide #2 C7.8-9 of the 216aa ORF of *C. trachomatis* serovar L2.

SEQ ID NO: 156 is the determined amino acid sequence for the peptide # 2 C7.8-10 of the 216aa ORF of *C. trachomatis* serovar L2.

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SEQ ID NO: 157 is the determined amino acid sequence for the 53 amino acid residue peptide of the 216aa ORF within clone 2C7.8 of *C. trachomatis* serovar L2.

SEQ ID NO: 158 is the determined amino acid sequence for the 52 amino acid residue peptide of the CT529 ORF within clone 2C7.8 of *C. trachomatis* serovar L2.

SEQ ID NO: 159 is the determined DNA sequence for the 5' (forward) primer for cloning full-length CT529 serovar L2.

SEQ ID NO: 160 is the determined DNA sequence for the 5' (reverse) primer for cloning full-length CT529 serovar L2.

SEQ ID NO: 161 is the determined DNA sequence for the 5' (forward) primer for cloning full-length CT529 for serovars other than L2 and MOPN.

SEQ ID NO: 162 is the determined DNA sequence for the 5' (reverse) primer for cloning full-length CT529 serovars other than L2 and MOPN.

SEQ ID NO: 163 is the determined DNA sequence for the 5' (forward) primer for cloning full-length CT529 serovar MOPN.

SEQ ID NO: 164 is the determined DNA sequence for the 5' (reverse) primer for cloning full-length CT529 serovar MOPN.

SEQ ID NO: 165 is the determined DNA sequence for the 5' (forward) primer for pBIB-KS.

SEQ ID NO: 166 is the determined DNA sequence for the 5' (reverse) primer for pBIB-KS.

SEQ ID NO: 167 is the determined amino acid sequence for the 9-mer epitope peptide Cap1#139-147 from serovar L2.

SEQ ID NO: 168 is the determined amino acid sequence for the 9-mer epitope peptide Cap1#139-147 from serovar D.

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SEQ ID NO: 169 is the determined full-length DNA sequence for the *C. trachomatis* pmpI (CT874) gene.

SEQ ID NO: 170 is the determined full-length DNA sequence for the *C. trachomatis* pmpG gene.

SEQ ID NO: 171 is the determined full-length DNA sequence for the *C. trachomatis* pmpE gene.

SEQ ID NO: 172 is the determined full-length DNA sequence for the *C. trachomatis* pmpD gene.

SEQ ID NO: 173 is the determined full-length DNA sequence for the C.

10 trachomatis pmpC gene.

SEQ ID NO: 174 is the determined full-length DNA sequence for the *C. trachomatis* pmpB gene.

SEQ ID NO: 175 is the predicted full-length amino acid sequence for the C. trachomatis pmpI gene.

SEQ ID NO: 176 is the predicted full-length amino acid sequence for the *C. trachomatis* pmpG gene.

SEQ ID NO: 177 is the predicted full-length amino acid sequence for the *C. trachomatis* pmpE gene.

SEQ ID NO: 178 is the predicted full-length amino acid sequence for the ...

C. trachomatis pmpD gene.

SEQ ID NO: 179 is the predicted full-length amino acid sequence for the *C. trachomatis* pmpC gene.

SEQ ID NO: 180 is the predicted full-length amino acid sequence for the *C. trachomatis* pmpB gene.

SEQ ID NO: 181 is the determined DNA sequence minus the signal sequence for the *C. trachomatis* pmpI gene.

SEQ ID NO: 182 is a subsequently determined full-length DNA sequence for the C. trachomatis pmpG gene.

SEQ ID NO: 183 is the determined DNA sequence minus the signal sequence for the C. trachomatis pmpE gene.

SEQ ID NO: 184 is a first determined DNA sequence representing the carboxy terminus for the C. trachomatis pmpD gene.

SEQ ID NO: 185 is a second determined DNA sequence representing the amino terminus minus the signal sequence for the *C. trachomatis* pmpD gene.

SEQ ID NO: 186 is a first determined DNA sequence representing the carboxy terminus for the *C. trachomatis* pmpC gene.

SEQ ID NO: 187 is a second determined DNA sequence representing the amino terminus minus the signal sequence for the *C. trachomatis* pmpC gene.

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SEQ ID NO: 188 is the determined DNA sequence representing the *C. pneumoniae* serovar MOMPS pmp gene in a fusion molecule with Ra12.

SEQ ID NO: 189 is the predicted amino acid sequence minus the signal sequence for the *C. trachomatis* pmpI gene.

SEQ ID NO: 190 is subsequently predicted amino acid sequence for the C. trachomatis pmpG gene.

SEQ ID NO: 191 is the predicted amino acid sequence minus the signal sequence for the *C. trachomatis* pmpE gene.

SEQ ID NO: 192 is a first predicted amino acid sequence representing the carboxy terminus for the *C. trachomatis* pmpD gene.

SEQ ID NO: 193 is a second predicted amino acid sequence representing the Amino terminus minus the signal sequence for the *C. trachomatis* pmpD gene.

SEQ ID NO: 194 is a first predicted amino acid sequence representing the Carboxy terminus for the *C. trachomatis* pmpC gene.

SEQ ID NO: 195 is a second predicted amino acid sequence representing the Amino terminus for the *C. trachomatis* pmpC gene.

SEQ ID NO: 196 is the predicted amino acid sequence representing the *C. pneumoniae* serovar MOMPS pmp gene in a fusion molecule with Ra12.

SEQ ID NO: 197 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpC gene in the SKB vaccine vector.

SEQ ID NO: 198 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpC gene in the SKB vaccine vector.

SEQ ID NO: 199 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpC gene in the SKB vaccine vector.

SEQ ID NO: 200 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpD gene in the SKB vaccine vector.

SEQ ID NO: 201 is the determined DNA sequence for the 3' oligo primer for cloning the C. trachomatis pmpD gene in the SKB vaccine vector.

SEQ ID NO: 202 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpD gene in the SKB vaccine vector.

SEQ ID NO: 203 is the determined DNA sequence for the 5' oligo primer for cloning the C. trachomatis pmpE gene in the SKB vaccine vector.

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SEQ ID NO: 204 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpE gene in the SKB vaccine vector.

SEQ ID NO: 205 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpG gene in the SKB vaccine vector.

SEQ ID NO: 206 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpG gene in the SKB vaccine vector.

SEQ ID NO: 207 is the determined DNA sequence for the 5' oligo primer for cloning the amino terminus portion of the *C. trachomatis* pmpC gene in the pET17b vector.

SEQ ID NO: 208 is the determined DNA sequence for the 3' oligo primer for cloning the amino terminus portion of the *C. trachomatis* pmpC gene in the pET17b vector.

SEQ ID NO: 209 is the determined DNA sequence for the 5' oligo primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpC gene in the pET17b vector.

SEQ ID NO: 210 is the determined DNA sequence for the 3' oligo primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpC gene in the pET17b vector.

SEQ ID NO: 211 is the determined DNA sequence for the 5' oligo primer for cloning the amino terminus portion of the *C. trachomatis* pmpD gene in the pET17b vector.

SEQ ID NO: 212 is the determined DNA sequence for the 3' oligo primer for cloning the amino terminus portion of the *C. trachomatis* pmpD gene in the pET17b vector.

SEQ ID NO: 213 is the determined DNA sequence for the 5' oligo primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpD gene in the pET17b vector.

SEQ ID NO: 214 is the determined DNA sequence for the 3' oligo primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpD gene in the pET17b vector.

SEQ ID NO: 215 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpE gene in the pET17b vector.

SEQ ID NO: 216 is the determined DNA sequence for the 3' oligo primer for cloning the C. trachomatis pmpE gene in the pET17b vector.

SEQ ID NO: 217 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpE gene in the pET17b vector.

SEQ ID NO: 218 is the amino acid sequence for the insertion sequence for cloning the *C. trachomatis* pmpE gene in the pET17b vector.

SEQ ID NO: 219 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpG gene in the pET17b vector.

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SEQ ID NO: 220 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpG gene in the pET17b vector.

SEQ ID NO: 221 is the amino acid sequence for the insertion sequence for cloning the *C. trachomatis* pmpG gene in the pET17b vector.

SEQ ID NO: 222 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpI gene in the pET17b vector.

SEQ ID NO: 223 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpI gene in the pET17b vector.

SEQ ID NO: 224 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 1-20.

SEQ ID NO: 225 is the determined amino acid sequence for the C. pneumoniae Swib peptide 6-25.

SEQ ID NO: 226 is the determined amino acid sequence for the C. 30 pneumoniae Swib peptide 12-31.

SEQ ID NO: 227 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 17-36.

SEQ ID NO: 228 is the determined amino acid sequence for the C. pneumoniae Swib peptide 22-41.

SEQ ID NO: 229 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 27-46.

5 SEQ ID NO: 230 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 42-61.

SEQ ID NO: 231 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 46-65.

SEQ ID NO: 232 is the determined amino acid sequence for the C.

10 pneumoniae Swib peptide 51-70.

SEQ ID NO: 233 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 56-75.

SEQ ID NO: 234 is the determined amino acid sequence for the C. pneumoniae Swib peptide 61-80.

SEQ ID NO: 235 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 66-87.

SEQ ID NO: 236 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 103-122.

SEQ ID NO: 237 is the determined amino acid sequence for the *C*.

20 trachomatis OMCB peptide 108-127.

SEQ ID NO: 238 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 113-132.

SEQ ID NO: 239 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 118-137.

SEQ ID NO: 240 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 123-143.

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SEQ ID NO: 241 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 128-147.

SEQ ID NO: 242 is the determined amino acid sequence for the *C.* trachomatis OMCB peptide 133-152.

SEQ ID NO: 243 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 137-156.

SEQ ID NO: 244 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 142-161.

SEQ ID NO: 245 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 147-166.

SEQ ID NO: 246 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 152-171.

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SEQ ID NO: 247 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 157-176.

SEQ ID NO: 248 is the determined amino acid sequence for the *C.*10 trachomatis OMCB peptide 162-181.

SEQ ID NO: 249 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 167-186.

SEQ ID NO: 250 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 171-190.

SEQ ID NO: 251 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 171-186.

SEQ ID NO: 252 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 175-186.

SEQ ID NO: 252 is the determined amino acid sequence for the *C.* 20 trachomatis OMCB peptide 175-186.

SEQ ID NO: 253 is the determined amino acid sequence for the *C. pneumoniae* OMCB peptide 185-198.

SEQ ID NO: 254 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 96-115.

SEQ ID NO: 255 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 101-120.

SEQ ID NO: 256 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 106-125.

SEQ ID NO: 257 is the determined amino acid sequence for the *C.* 30 trachomatis TSA peptide 111-130.

SEQ ID NO: 258 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 116-135.

SEQ ID NO: 259 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 121-140.

SEQ ID NO: 260 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 126-145.

SEQ ID NO: 261 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 131-150.

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SEQ ID NO: 262 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 136-155.

SEQ ID NO: 263 is the determined full-length DNA sequence for the C.

trachomatis CT529/Cap 1 gene serovar I.

SEQ ID NO: 264 is the predicted full-length amino sequence for the *C. trachomatis* CT529/Cap 1 gene serovar I.

SEQ ID NO: 265 is the determined full-length DNA sequence for the C. trachomatis CT529/Cap 1 gene serovar K.

SEQ ID NO: 266 is the predicted full-length amino sequence for the *C. trachomatis* CT529/Cap 1 gene serovar K.

SEQ ID NO: 267 is the determined DNA sequence for the *C. trachomatis* clone 17-G4-36 sharing homology to part of the ORF of DNA-dirrected RNA polymerase beta subunit- CT315 in serD.

SEQ ID NO: 268 is the determined DNA sequence for the partial sequence of the *C. trachomatis* CT016 gene in clone 2E10.

SEQ ID NO: 269 is the determined DNA sequence for the partial sequence of the *C. trachomatis* tRNA syntase gene in clone 2E10.

SEQ ID NO: 270 is the determined DNA sequence for the partial sequence for the *C. trachomatis* clpX gene in clone 2E10.

SEQ ID NO: 271 is a first determined DNA sequence for the *C. trachomatis* clone CtL2gam-30 representing the 5'end.

SEQ ID NO: 272 is a second determined DNA sequence for the *C. trachomatis* clone CtL2gam-30 representing the 3'end.

SEQ ID NO: 273 is the determined DNA sequence for the C. trachomatis clone CtL2gam-28.

SEQ ID NO: 274 is the determined DNA sequence for the *C. trachomatis* clone CtL2gam-27.

SEQ ID NO: 275 is the determined DNA sequence for the *C. trachomatis* clone CtL2gam-26.

SEQ ID NO: 276 is the determined DNA sequence for the C. trachomatis clone CtL2gam-24.

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SEQ ID NO: 277 is the determined DNA sequence for the *C. trachomatis* clone CtL2gam-23.

SEQ ID NO: 278 is the determined DNA sequence for the C. 10 trachomatis clone CtL2gam-21.

SEQ ID NO: 279 is the determined DNA sequence for the *C. trachomatis* clone CtL2gam-18.

SEQ ID NO: 280 is the determined DNA sequence for the C. trachomatis clone CtL2gam-17.

SEQ ID NO: 281 is a first determined DNA sequence for the C. trachomatis clone CtL2gam-15 representing the 5' end.

SEQ ID NO: 282 is a second determined DNA sequence for the *C. trachomatis* clone CtL2gam-15 representing the 3' end.

SEQ ID NO: 283 is the determined DNA sequence for the C. 20 trachomatis clone CtL2gam-13.

SEQ ID NO: 284 is the determined DNA sequence for the *C. trachomatis* clone CtL2gam-10.

SEQ ID NO: 285 is the determined DNA sequence for the C. trachomatis clone CtL2gam-8.

SEQ ID NO: 286 is a first determined DNA sequence for the C. trachomatis clone CtL2gam-6 representing the 5' end.

SEQ ID NO: 287 is a second determined DNA sequence for the C. trachomatis clone CtL2gam-6 representing the 3' end.

SEQ ID NO: 288 is the determined DNA sequence for the C. 30 trachomatis clone CtL2gam-5.

SEQ ID NO: 289 is the determined DNA sequence for the C. trachomatis clone CtL2gam-2.

SEQ ID NO: 290 is the determined DNA sequence for the *C. trachomatis* clone CtL2gam-1.

SEQ ID NO: 291 is the determined full-length DNA sequence for the *C. pneumoniae* homologue of the CT529 gene.

SEQ ID NO: 292 is the predicted full-length amino acid sequence for the *C. pneumoniae* homologue of the CT529 gene.

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SEQ ID NO: 293 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpG gene in the SKB vaccine vector.

SEQ ID NO: 294 is the amino acid sequence of an open reading frame of clone CT603.

SEQ ID NO: 295 is the amino acid sequence of a first open reading frame of clone CT875.

SEQ ID NO: 296 is the amino acid sequence of a second open reading frame of clone CT875.

SEQ ID NO: 297 is the amino acid sequence of a first open reading frame of clone CT858.

SEQ ID NO: 298 is the amino acid sequence of a second open reading frame of clone CT858.

SEQ ID NO: 299 is the amino acid sequence of an open reading frame of clone CT622.

SEQ ID NO: 300 is the amino acid sequence of an open reading frame of clone CT610.

SEQ ID NO: 301 is the amino acid sequence of an open reading frame of clone CT396.

SEQ ID NO: 302 is the amino acid sequence of an open reading frame of clone CT318.

SEQ ID NO: 304 is the amino acid sequence for *C. trachomatis*, serovar L2 rCt529c1-125 having a modified N-terminal sequence (6-His tag).

SEQ ID NO: 305 is the amino acid sequence for *C. trachomatis*, serovar L2 rCt529c1-125.

SEQ ID NO: 306 is the sense primer used in the synthesis of the PmpA(N-term) fusion protein.

SEQ ID NO: 307 is the antisense primer used in the synthesis of the PmpA(N-term) fusion protein.

SEQ ID NO: 308 is the DNA sequence encoding the PmpA(N-term) fusion protein.

5 SEQ ID NO: 309 is the amino acid sequence of the PmpA(N-term) fusion protein.

SEQ ID NO: 310 is the sense primer used in the synthesis of the PmpA(C-term) fusion protein.

SEQ ID NO: 311 is the antisense primer used in the synthesis of the PmpA(C-term) fusion protein.

SEQ ID NO: 312 is the DNA sequence encoding the PmpA(C-term) fusion protein.

SEQ ID NO: 313 is the amino acid sequence of the PmpA(C-term) fusion protein.

SEQ ID NO: 314 is the sense primer used in the synthesis of the PmpF(N-term) fusion protein.

SEQ ID NO: 315 is the antisense primer used in the synthesis of the PmpF(N-term) fusion protein.

SEQ ID NO: 316 is the DNA sequence encoding the PmpF(N-term) fusion protein.

SEQ ID NO: 317 is the amino acid sequence of the PmpF(N-term) fusion protein.

SEQ ID NO: 318 is the sense primer used in the synthesis of the PmpF(C-term) fusion protein.

SEQ ID NO: 319 is the antisense primer used in the synthesis of the PmpF(C-term) fusion protein.

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SEQ ID NO: 320 is the DNA sequence encoding the PmpF(C-term) fusion protein.

SEQ ID NO: 321 is the amino acid sequence of the PmpF(C-term) fusion protein.

SEQ ID NO: 322 is the sense primer used in the synthesis of the PmpH (CT412) (N-term) fusion protein.

SEQ ID NO: 323 is the antisense primer used in the synthesis of the PmpH(N-term) fusion protein.

SEQ ID NO: 324 is the DNA sequence encoding the PmpH(N-term) fusion protein.

5 SEQ ID NO: 325 is the amino acid sequence of the PmpH(N-term) fusion protein.

SEQ ID NO: 326 is the sense primer used in the synthesis of the PmpH(C-term) fusion protein.

SEQ ID NO: 327 is the antisense primer used in the synthesis of the PmpH(C-term) fusion protein.

SEQ ID NO: 328 is the DNA sequence encoding the PmpH(C-term) fusion protein.

SEQ ID NO: 329 is the amino acid sequence of the PmpH(C-term) fusion protein.

SEQ ID NO: 330 is the sense primer used in the synthesis of the PmpB(1) fusion protein.

SEQ ID NO: 331 is the antisense primer used in the synthesis of the PmpB(1) fusion protein.

SEQ ID NO: 332 is the DNA sequence encoding the PmpB(1) fusion 20 protein.

SEQ ID NO: 333 is the amino acid sequence of the PmpB(1) fusion protein.

SEQ ID NO: 334 is the sense primer used in the synthesis of the PmpB(2) fusion protein.

SEQ ID NO: 335 is the antisense primer used in the synthesis of the PmpB(2) fusion protein.

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SEQ ID NO: 336 is the DNA sequence encoding the PmpB(2) fusion protein.

SEQ ID NO: 337 is the amino acid sequence of the PmpB(2) fusion 30 protein.

SEQ ID NO: 338 is the sense primer used in the synthesis of the PmpB(3) fusion protein.

SEQ ID NO: 339 is the antisense primer used in the synthesis of the PmpB(3) fusion protein.

SEQ ID NO: 340 is the DNA sequence encoding the PmpB(3) fusion protein.

SEQ ID NO: 341 is the amino acid sequence of the PmpB(3) fusion protein.

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SEQ ID NO: 342 is the sense primer used in the synthesis of the PmpB(4) fusion protein.

SEQ ID NO: 343 is the antisense primer used in the synthesis of the 10 PmpB(4) fusion protein.

SEQ ID NO: 344 is the DNA sequence encoding the PmpB(4) fusion protein.

SEQ ID NO: 345 is the amino acid sequence of the PmpB(4) fusion protein.

SEQ ID NO: 346 is the sense primer used in the synthesis of the PmpC(1) fusion protein.

SEQ ID NO: 347 is the antisense primer used in the synthesis of the PmpC(1) fusion protein.

SEQ ID NO: 348 is the DNA sequence encoding the PmpC(1) fusion 20 protein.

SEQ ID NO: 349 is the amino acid sequence of the PmpC(1) fusion protein.

SEQ ID NO: 350 is the sense primer used in the synthesis of the PmpC(2) fusion protein.

SEQ ID NO: 351 is the antisense primer used in the synthesis of the PmpC(2) fusion protein.

SEQ ID NO: 352 is the DNA sequence encoding the PmpC(2) fusion protein.

SEQ ID NO: 353 is the amino acid sequence of the PmpC(2) fusion 30 protein.

SEQ ID NO: 354 is the sense primer used in the synthesis of the PmpC(3) fusion protein.

SEQ ID NO: 355 is the antisense primer used in the synthesis of the PmpC(3) fusion protein.

SEQ ID NO: 356 is the DNA sequence encoding the PmpC(3) fusion protein.

SEQ ID NO: 357 is the amino acid sequence of the PmpC(3) fusion protein.

SEQ ID NO: 358 is the DNA sequence of the oppA1 protein, devoid of the first trans-membrane domain.

SEQ ID NO: 359 is the full length DNA sequence of CT139.

SEQ ID NO: 360 is the full length DNA sequence of ORF-3.

SEQ ID NO: 361 is the full length DNA sequence of CT611.

SEQ ID NO: 362 is the amino acid sequence of oppA1 starting from amino acid 22.

SEQ ID NO: 363 is the amino acid sequence of CT139.

SEQ ID NO: 364 is the amino acid sequence of ORF-3.

SEQ ID NO: 365 is the amino acid sequence of CT611.

SEQ ID NO: 366 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0275, of the Chlamydia trachomatis gene CT190.

SEQ ID NO: 367 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0407, of the Chlamydia trachomatis gene CT103.

SEQ ID NO: 368 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0720, of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 369 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0716, of the Chlamydia trachomatis gene CT660.

SEQ ID NO: 370 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0519, of the Chlamydia trachomatis gene CT430.

SEQ ID NO: 371 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0520, of the Chlamydia trachomatis gene CT431.

SEQ ID NO: 372 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0078, of the Chlamydia trachomatis gene CT318.

SEQ ID NO: 373 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0628, of the Chlamydia trachomatis gene CT509.

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SEQ ID NO: 374 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0540, of the Chlamydia trachomatis gene CT414.

SEQ ID NO: 375 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, pmp20, of the Chlamydia trachomatis gene CT413.

SEQ ID NO: 376 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0081, of the Chlamydia trachomatis gene CT315.

SEQ ID NO: 377 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0761, of the Chlamydia trachomatis gene CT610.

SEQ ID NO: 378 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0557, of the Chlamydia trachomatis gene CT443.

SEQ ID NO: 379 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0833, of the Chlamydia trachomatis gene CT557.

SEQ ID NO: 380 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0134, of the Chlamydia trachomatis gene CT604.

SEQ ID NO: 381 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0388, of the Chlamydia trachomatis gene CT042.

SEQ ID NO: 382 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn1028, of the Chlamydia trachomatis gene CT376.

SEQ ID NO: 383 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0875, of the Chlamydia trachomatis gene CT734.

SEQ ID NO: 384 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0908, of the Chlamydia trachomatis gene CT764.

SEQ ID NO: 385 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0728, of the Chlamydia trachomatis gene CT622.

SEQ ID NO: 386 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0275, of the Chlamydia trachomatis gene CT190.

SEQ ID NO: 387 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0407, of the Chlamydia trachomatis gene CT103.

SEQ ID NO: 388 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0720, of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 389 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0716, of the Chlamydia trachomatis gene CT660.

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SEQ ID NO: 390 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0519, of the Chlamydia trachomatis gene CT430. SEQ ID NO: 391 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0520, of the Chlamydia trachomatis gene CT431. SEO ID NO: 392 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0078, of the Chlamydia trachomatis gene CT318. SEO ID NO: 393 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0628, of the Chlamydia trachomatis gene CT509. SEQ ID NO: 394 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0540, of the Chlamydia trachomatis gene CT414. SEO ID NO: 395 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, pmp20, of the Chlamydia trachomatis gene CT413. SEQ ID NO: 396 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0081, of the Chlamydia trachomatis gene CT315. SEQ ID NO: 397 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0761, of the Chlamydia trachomatis gene CT610. SEQ ID NO: 398 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0557, of the Chlamydia trachomatis gene CT443. SEQ ID NO: 399 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0833, of the Chlamydia trachomatis gene CT557. SEQ ID NO: 400 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0134, of the Chlamydia trachomatis gene CT604. SEQ ID NO: 401 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0388, of the Chlamydia trachomatis gene CT042. SEQ ID NO: 402 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn1028, of the Chlamydia trachomatis gene CT376. SEQ ID NO: 403 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0875, of the Chlamydia trachomatis gene CT734. SEQ ID NO: 404 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0908, of the Chlamydia trachomatis gene CT764. SEQ ID NO: 405 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0728, of the Chlamydia trachomatis gene CT622.

SEQ ID NO: 406 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT287.

SEQ ID NO: 407 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT858.

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SEQ ID NO: 408 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT764.

SEQ ID NO: 409 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT734.

SEQ ID NO: 410 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT660.

SEQ ID NO: 411 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 412 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT622.

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SEQ ID NO: 413 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT610.

SEQ ID NO: 414 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT604.

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SEQ ID NO: 415 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT557.

SEQ ID NO: 416 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT509.

SEQ ID NO: 417 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT443.

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SEQ ID NO: 418 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT431.

SEQ ID NO: 419 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT430.

SEQ ID NO: 420 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT414.

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SEQ ID NO: 421 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT413.

SEQ ID NO: 422 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT396.

SEQ ID NO: 423 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT376.

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SEQ ID NO: 424 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT318.

SEQ ID NO: 425 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT315.

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SEQ ID NO: 426 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT104.

SEQ ID NO: 427 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT103.

SEQ ID NO: 428 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT102.

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SEQ ID NO: 429 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT098.

SEQ ID NO: 430 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT042.

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SEQ ID NO: 431 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT858.

SEQ ID NO: 432 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT764.

SEQ ID NO: 433 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT734.

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SEQ ID NO: 434 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT660.

SEQ ID NO: 435 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 436 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT622.

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SEQ ID NO: 437 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT610.

SEQ ID NO: 438 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT604.

SEQ ID NO: 439 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT557.

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SEQ ID NO: 440 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT509.

SEQ ID NO: 441 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT443.

SEQ ID NO: 442 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT431.

SEQ ID NO: 443 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT430.

SEQ ID NO: 444 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT414.

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SEQ ID NO: 445 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT413.

SEQ ID NO: 446 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT396.

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SEQ ID NO: 447 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT376.

SEQ ID NO: 448 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT318.

SEQ ID NO: 449 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT315.

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SEQ ID NO: 450 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT104.

SEQ ID NO: 451 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT103.

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SEQ ID NO: 452 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT102.

SEQ ID NO: 453 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT098.

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SEQ ID NO: 454 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT042.

SEQ ID NO: 455 corresponds to the DNA sequence of CPn0894, which is the CP homologue of CT751 (amn), which was identified in clones CTL2-1, and CTL2-5.

SEQ ID NO: 456 corresponds to the DNA sequence of CPn0074, which is the CP homologue of CT322 (tuf), which was identified in clone CTL2-2.

SEQ ID NO: 457 corresponds to the DNA sequence of CPn0122, which is the CP homologue of CT032 (metG), which was identified in clones CTL2gam2, CTL2-3(5') and CTL2-4.

SEQ ID NO: 458 corresponds to the DNA sequence of CPn0121, which is the CP homologue of CT031, which was identified in clone CTL2-3(5')(3').

SEQ ID NO: 459 corresponds to the DNA sequence of CPn0120, which is the CP homologue of CT030 (gmK), which was identified in clones CTL2-3(3') and CTL2-21.

SEQ ID NO: 460 corresponds to the DNA sequence of CPn0359, which is the CP homologue of CT064 (lepA), which was identified in clone CTL2gam5.

SEQ ID NO: 461 corresponds to the DNA sequence of CPn0414, which is the CP homologue of CT265 (accA), which was identified in clone CTL2-6.

SEQ ID NO: 462 corresponds to the DNA sequence of CPn0413, which is the CP homologue of CT264 (msbA), which was identified in clone CTL2-6.

SEQ ID NO: 463 corresponds to the DNA sequence of CPn0394, which is the CP homologue of CT256 which was identified in clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 464 corresponds to the DNA sequence of CPn0395, which is the CP homologue of CT257 which was identified in clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 465 corresponds to the DNA sequence of CPn0487, which is the CP homologue of CT384 which was identified in clones CTL2gam6(3') and CTL2-11(3').

PCT/US01/23121

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SEQ ID NO: 466 corresponds to the DNA sequence of CPn0592, which is the CP homologue of CT473, which was identified in clone CTL2-8b.

SEQ ID NO: 467 corresponds to the DNA sequence of CPn0593, which is the CP homologue of CT474, which was identified in clone CTL2-8b.

SEQ ID NO: 468 corresponds to the DNA sequence of CPn0197, which is the CP homologue of CT139 (oppA1), which was identified in clone CTL2-8b.

SEQ ID NO: 469 corresponds to the DNA sequence of CPn0363, which is the CP homologue of CT060 (flhA), which was identified in clone CTL2-8b.

SEQ ID NO: 470 corresponds to the DNA sequence of CPn0301, which is the CP homologue of CT242, which was identified in clone CTL2gam8.

SEQ ID NO: 471 corresponds to the DNA sequence of CPn0302, which is the CP homologue of CT243 (lpxD), which was identified in clone CTL2gam8.

SEQ ID NO: 472 corresponds to the DNA sequence of CPn0324, which is the CP homologue of CT089 (lcrE), which was identified in clones CTL2-9, CTL2gam1, CTL2gam17 and CTL2-19(5').

SEQ ID NO: 473 corresponds to the DNA sequence of CPn0761, which is the CP homologue of CT610, which was identified in clone CTL2-10(5')(3').

SEQ ID NO: 474 corresponds to the DNA sequence of CPn0760, which is the CP homologue of CT611, which was identified in clone CTL2-10(5').

SEQ ID NO: 475 corresponds to the DNA sequence of CPn0329, which is the CP homologue of CT154, which was identified in clones CTL2gam10 and CTL2gam21.

SEQ ID NO: 476 corresponds to the DNA sequence of CPn0990, which is the CP homologue of CT833 (infC), which was identified in clone CTL2-12.

SEQ ID NO: 477 corresponds to the DNA sequence of CPn0984, which is the CP homologue of CT827 (nrdA), which was identified in clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 478 corresponds to the DNA sequence of CPn0985 which is the CP homologue of CT828 (nrdB) which was identified in clones CTL2-16(3') CTL2gam15(3').

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WO 02/08267

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SEQ ID NO: 479 corresponds to the DNA sequence of CPn0349, which is the CP homologue of CT067 (ytgA), which was identified in clone CTL2gam18.

SEQ ID NO: 480 corresponds to the DNA sequence of CPn0325, which is the CP homologue of CT088 (sycE), which was identified in clone CTL2-19(5').

SEQ ID NO: 481 corresponds to the DNA sequence of CPn0326, which is the CP homologue of CT087 (malQ), which was identified in clone CTL2-19(5').

SEQ ID NO: 482 corresponds to the DNA sequence of CPn0793, which is the CP homologue of CT588 (rbsu), which was identified in clone CTL2gam23.

SEQ ID NO: 483 corresponds to the DNA sequence of CPn0199, which is the CP homologue of CT199 (oppB1), which was identified in clone CTL2gam24.

SEQ ID NO: 484 corresponds to the DNA sequence of CPn0666, which is the CP homologue of CT545 (dnaE), which was identified in clone CTL2-24.

SEQ ID NO: 485 corresponds to the DNA sequence of CPn0065, which is the CP homologue of CT288, which was identified in clone CTL2gam27.

SEQ ID NO: 486 corresponds to the DNA sequence of CPn0444, which is the CP homologue of CT413 (pmpB), which was identified in clone CTL2gam30(5')(3').

SEQ ID NO: 487 corresponds to the DNA sequence of CPn-ORF5, which is the CP homologue of CT-ORF3, which was identified in clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5'), and CTL2-23.

SEQ ID NO: 488 corresponds to the DNA sequence of CPn-ORF6, which is the CP homologue of CT-ORF4, which was identified in clone CTL2-18(3').

SEQ ID NO: 489 corresponds to the DNA sequence of CP-ORF7, which is the CP homologue of CT-ORF5, which was identified in clone CTL2-18(3').

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SEQ ID NO: 490 corresponds to the amino acid sequence of CPn0894, which is the CP homologue of CT751 (amn), which was identified in clones CTL2-1 and CTL2-5.

SEQ ID NO: 491 corresponds to the amino acid sequence of CPn0074, which is the CP homologue of CT332 (tuf), which was identified in clone CTL2-2.

SEQ ID NO: 492 corresponds to the amino acid sequence of CPn0122, which is the CP homologue of CT032 (metG), which was identified in clones CTL2gam2, CTL2-3(5') and CTL2-4.

SEQ ID NO: 493 corresponds to the amino acid sequence of CPn0121, which is the CP homologue of CT031, which was identified in clone CTL2-3(5')(3').

SEQ ID NO: 494 corresponds to the amino acid sequence of CPn0120 which is the CP homologue of CT030 (gmK) which was identified in clones CTL2-3 (3') and CTL2-21.

SEQ ID NO: 495 corresponds to the amino acid sequence of CPn0359, which is the CP homologue of CT064 (lepA), which was identified in clone CTL2gam5.

SEQ ID NO: 496 corresponds to the amino acid sequence of CPn0414, which is the CP homologue of CT265 (accA), which was identified in clone CTL2-6.

SEQ ID NO: 497 corresponds to the amino acid sequence of CPn0413, which is the CP homologue of CT264 (msbA), which was identified in clone CTL2-6.

SEQ ID NO: 498 corresponds to the amino acid sequence of CPn0394, which is the CP homologue of CT256, which was identified in clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 499 corresponds to the amino acid sequence of CPn0395, which is the CP homologue of CT257, which was identified in clones CTL2gam6(5') and CTL2-11(5').

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WO 02/08267 PCT/US01/23121

SEQ ID NO: 500 corresponds to the amino acid sequence of CPn0487, which is the CP homologue of CT384, which was identified in clones CTL2gam6(3') and CTL2-11(3').

SEQ ID NO: 501 corresponds to the amino acid sequence of CPn0592, which is the CP homologue of CT473, which was identified in clone CTL2-8b.

SEQ ID NO: 502 corresponds to the amino acid sequence of CPn0593, which is the CP homologue of CT474, which was identified in clone CTL2-8b.

SEQ ID NO: 503 corresponds to the amino acid sequence of CPn0197, which is the CP homologue of CT139 (oppA1), which was identified in clone CTL2-8b.

SEQ ID NO: 504 corresponds to the amino acid sequence of CPn0363, which is the CP homologue of CT060 (flhA), which was identified in clone CTL2-8b.

SEQ ID NO: 505 corresponds to the amino acid sequence of CPn0301, which is the CP homologue of CT242, which was identified in clone CTL2gam8.

SEQ ID NO: 506 corresponds to the amino acid sequence of CPn0302, which is the CP homologue of CT243 (lpxD), which was identified in clone CTL2gam8.

SEQ ID NO: 507 corresponds to the amino acid sequence of CPn0324, which is the CP homologue of CT089 (lcrE), which was identified in clones CTL2-9, CTL2gam1, CTL2gam17 and CTL2-19(5').

SEQ ID NO: 508 corresponds to the amino acid sequence of CPn0761, which is the CP homologue of CT610, which was identified in clone CTL2-10(5')(3').

SEQ ID NO: 509 corresponds to the amino acid sequence of CPn0760, which is the CP homologue of CT611, which was identified in clone CTL2-10(5').

SEQ ID NO: 510 corresponds to the amino acid sequence of CPn0329, which is the CP homologue of CT154, which was identified in clones CTL2gam10 and CTL2gam21.

SEQ ID NO: 511 corresponds to the amino acid sequence of CPn0990, which is the CP homologue of CT833 (infC), which was identified in clone CTL2-12.

SEQ ID NO: 512 corresponds to the amino acid sequence of CPn-ORF5, which is the CP homologue of CT ORF3, which was identified in clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5'), and CTL2-23.

SEQ ID NO: 513 corresponds to the amino acid sequence of CPn0984, which is the CP homologue of CT827 (nrdA) which was identified in clones CTL2-16(3') and CTL2gam15(3').

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SEQ ID NO: 514 corresponds to the amino acid sequence of CPn0985, which is the CP homologue of CT828 (nrdB) which was identified in clones CTL2-16(3') CTL2gam15(3').

SEQ ID NO: 515 corresponds to the amino acid sequence of CPn0349, which is the CP homologue of CT067 (ytgA), which was identified in clone CTL2gam18.

SEQ ID NO: 516 corresponds to the DNA sequence of CPn-ORF6, which is the CP homologue of CT-ORF4, which was identified in clone CTL2-18(3').

SEQ ID NO: 517 corresponds to the DNA sequence of CP-ORF7, which is the CP homologue of CT-ORF5, which was identified in clone CTL2-18(3').

SEQ ID NO: 518 corresponds to the amino acid sequence of CPn0326, which is the CP homologue of CT087 (malQ), which was identified in clone CTL2-19(5').

SEQ ID NO: 519 corresponds to the amino acid sequence of CPn0325, which is the CP homologue of CT088 (sycE), which was identified in clone CTL2-19(5').

SEQ ID NO: 520 corresponds to the amino acid sequence of CPn0793, which is the CP homologue of CT588 (rbsu), which was identified in clone CTL2gam23.

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SEQ ID NO: 521 corresponds to the amino acid sequence of CPn0199, which is the CP homologue of CT199 (oppB1), which was identified in clone CTL2gam24.

SEQ ID NO: 522 corresponds to the amino acid sequence of CPn0666, which is the CP homologue of CT545 (dnaE), which was identified in clone CTL2-24.

SEQ ID NO: 523 corresponds to the DNA sequence of CPn0065, which is the CP homologue of CT288, which was identified in clone CTL2gam27.

SEQ ID NO: 524 corresponds to the DNA sequence of CPn0444, which is the CP homologue of CT413 (pmpB), which was identified in clone CTL2gam30(5')(3').

SEQ ID NO: 525 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT751 (amn) identified from the clones CTL2-1 and CTL2-5.

SEQ ID NO: 526 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT322 (tuff) identified from the clone CTL2-2.

SEQ ID NO: 527 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT032 (metG) identified from the clones CTL2gam2, CTL2-3(5') and CTL2-4.

SEQ ID NO: 528 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT031 identified from the clone CTL2-3(5')(3').

SEQ ID NO: 529 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT030 (gmK) identified from the clones CTL2-3(3') and CTL2-21.

SEQ ID NO: 530 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT064 (lepA) identified from the clone CTL2gam5.

SEQ ID NO: 531 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT265 (accA) identified from the clone CTL2-6.

SEQ ID NO: 532 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT624 (msbA) identified from the clones CTL2-6.

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SEQ ID NO: 533 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT256 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 534 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT257 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 535 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT384 identified from the clones CTL2gam6(3') and CTL2-11(3').

SEQ ID NO: 536 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT473 identified from the clone CTL2-8b.

SEQ ID NO: 537 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT474 identified from the clones CTL2-8b.

SEQ ID NO: 538 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT139 (oppA1) identified from the clones CTL2-8b.

SEQ ID NO: 539 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT060 (flhA) identified from the clone CTL2-8b.

SEQ ID NO: 540 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT242 identified from the clone CTL2gam8.

SEQ ID NO: 541 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT243 (lpxD) identified from the clone CTL2gam8.

SEQ ID NO: 542 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT089 identified from the clones CTL2-9, CTL2gam1, CTL2gam17, and CTL2-19(5').

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SEQ ID NO: 543 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT610 identified from the clone CTL2-10 (5')(3').

SEQ ID NO: 544 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT611 identified from the clone CTL2-10(5').

SEQ ID NO: 545 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT154 identified from the clones CTL2gam10 and CTL2gam21.

SEQ ID NO: 546 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT833 (infC) identified from the clone CTL2-12.

SEQ ID NO: 547 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT827 (nrdA) identified from the clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 548 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT828 (nrdB) identified from the clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 549 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT067 (ytgA) identified from the clone CTL2gam18.

SEQ ID NO: 550 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT088 (sycE) identified from the clones CTL2-19(5').

SEQ ID NO: 551 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT087 identified from the clone CTL2-19(5').

SEQ ID NO: 552 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT588 (rsbu) identified from the clone CTL2gam23.

SEQ ID NO: 553 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT199 (oppB1) identified from the clone CTL2gam24.

SEQ ID NO: 554 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT545 (dnaE) identified from the clone CTL2-4.

SEQ ID NO: 555 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT288 identified from the clones CTL2gam27.

SEQ ID NO: 556 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT413 (pmpB) identified from the clone CTL2gam30(5')(3').

SEQ ID NO: 557 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT-ORF3 identified from the clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5') and CTL2-23.

SEQ ID NO: 558 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for pCT-ORF4 identified from the clone CTL2-18(3').

SEQ ID NO: 559 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT-ORF5 identified from the clones CTL2-18(3').

SEQ ID NO: 560 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT751 (amn) identified from the clones CTL2-1 and CTL2-5.

SEQ ID NO: 561 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT322 (tuff) identified from the clone CTL2-2.

SEQ ID NO: 562 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT032 (metG) identified from the clones CTL2gam2, CTL2-3(5') and CTL2-4.

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SEQ ID NO: 563 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT031 identified from the clone CTL2-3(5')(3').

SEQ ID NO: 564 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT030 (gmK) identified from the clones CTL2-3(3') and CTL2-21.

SEQ ID NO: 565 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT064 (lepA) identified from the clone CTL2gam5.

SEQ ID NO: 566 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT265 (accA) identified from the clone CTL2-6.

SEQ ID NO: 567 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT624 (msbA) identified from the clones CTL2-6.

SEQ ID NO: 568 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT256 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 569 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT257 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 570 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT384 identified from the clones CTL2gam6(3') and CTL2-11(3').

SEQ ID NO: 571 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT473 identified from the clone CTL2-8b.

SEQ ID NO: 572 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT474 identified from the clones CTL2-8b.

WO 02/08267

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SEQ ID NO: 573 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT139 (oppA1) identified from the clones CTL2-8b.

PCT/US01/23121

SEQ ID NO: 574 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT060 (flhA) identified from the clone CTL2-8b.

SEQ ID NO: 575 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT242 identified from the clone CTL2gam8.

SEQ ID NO: 576 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT243 (lpxD) identified from the clone CTL2gam8.

SEQ ID NO: 577 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT089 identified from the clones CTL2-9, CTL2gam1, CTL2gam17, and CTL2-19(5').

SEQ ID NO: 578 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT610 identified from the clone CTL2-10 (5')(3').

SEQ ID NO: 579 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT611 identified from the clone CTL2-10(5').

SEQ ID NO: 580 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT154 identified from the clones CTL2gam10 and CTL2gam21.

SEQ ID NO: 581 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT833 (infC) identified from the clone CTL2-12.

SEQ ID NO: 582 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT-ORF3 identified from the clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5') and CTL2-23.

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SEQ ID NO: 583 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT827 (nrdA) identified from the clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 584 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT828 (nrdB) identified from the clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 585 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT067 (ytgA) identified from the clone CTL2gam18.

SEQ ID NO: 586 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for pCT-ORF4 identified from the clone CTL2-18(3')

SEQ ID NO: 587 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT-ORF5 identified from the clones CTL2-18(3').

SEQ ID NO: 588 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT087 identified from the clone CTL2-19(5').

SEQ ID NO: 589 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT088 (sycE) identified from the clones CTL2-19(5').

SEQ ID NO: 590 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT588 (rsbu) identified from the clone CTL2gam23.

SEQ ID NO: 591 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT199 (oppB1) identified from the clone CTL2gam24.

SEQ ID NO: 592 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT545 (dnaE) identified from the clone CTL2-4.

SEQ ID NO: 593 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT288 identified from the clones CTL2gam27.

SEQ ID NO: 594 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT413 (pmpB) identified from the clone CTL2gam30(5')(3').

SEQ ID NO: 595 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0406, of the Chlamydia trachomatis gene CT102.

SEQ ID NO: 596 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0315, of the Chlamydia trachomatis gene CT098.

SEQ ID NO: 597 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0406, of the Chlamydia trachomatis gene CT102.

SEQ ID NO: 598 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0315, of the Chlamydia trachomatis gene CT098.

SEQ ID NO: 599 sets forth the amino acid sequence for Chlamydia trachomatis serovar D CT287 protein.

DESCRIPTION OF THE FIGURES

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- Fig. 1 illustrates induction of INF-γ from a *Chlamydia*-specific T cell line activated by target cells expressing clone 4C9-18#2.
- Fig. 2 illustrates retroviral vectors pBIB-KS1,2,3 modified to contain a Kosak translation initiation site and stop codons.
- Fig. 3 shows specific lysis in a chromium release assay of P815 cells pulsed with *Chlamydia* peptides CtC7.8-12 (SEQ ID NO: 18) and CtC7.8-13 (SEQ ID NO: 19).
- Fig. 4 shows antibody isotype titers in C57Bl/6 mice immunized with *C. trachomatis* SWIB protein.
 - Fig. 5 shows *Chlamydia*-specific T-cell proliferative responses in splenocytes from C3H mice immunized with *C. trachomatis* SWIB protein.
- Fig. 6 illustrates the 5' and 3' primer sequences designed from *C. pneumoniae* which were used to isolate the SWIB and S13 genes from *C. pneumoniae*.

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Figs. 7A and 7B show induction of IFN-γ from a human anti-chlamydia T-cell line (TCL-8) capable of cross-reacting to C. trachomatis and C. pneumonia upon activation by monocyte-derived dendritic cells expressing chlamydial proteins.

Fig. 8 shows the identification of T cell epitopes in Chlamydial ribosomal S13 protein with T-cell line TCL 8 EB/DC.

Fig. 9A and B illustrate the proliferative response of CP-21 T-cells generated against *C. pnuemoniae*-infected dendritic cells to recombinant *C. pneumonia*-SWIB protein, but not *C. trachomatis* SWIB protein.

Fig. 10 shows the *C. trachomatis*-specific SWIB proliferative responses of a primary T-cell line (TCT-10 EB) from an asymptomatic donor.

Fig. 11 illustrates the identification of T-cell epitope in *C. trachomatis* SWIB with an antigen specific T-cell line (TCL-10 EB).

Fig. 12 shows the *C. trachomatis*-specific proliferative responses of primary T cell lines generated from two patients against the CT specific antigens CT622, CT875 and CT EB.

DETAILED DESCRIPTION OF THE INVENTION

As noted above, the present invention is generally directed to compositions and methods for the diagnosis and treatment of Chlamydial infection. In one aspect, the compositions of the subject invention include polypeptides that comprise at least one immunogenic portion of a *Chlamydia* antigen, or a variant thereof.

In specific embodiments, the subject invention discloses polypeptides comprising an immunogenic portion of a *Chlamydia* antigen, wherein the *Chlamydia* antigen comprises an amino acid sequence encoded by a polynucleotide molecule disclosed herein, the complements of said nucleotide sequences, and variants of such sequences.

As used herein, the term "polypeptide" encompasses amino acid chains of any length, including full length proteins (i.e., antigens), wherein the amino acid residues are linked by covalent peptide bonds. Thus, a polypeptide comprising an immunogenic portion of one of the inventive antigens may consist entirely of the immunogenic portion, or may contain additional sequences. The additional sequences

49

may be derived from the native *Chlamydia* antigen or may be heterologous, and such sequences may (but need not) be immunogenic.

The term "polynucleotide(s)," as used herein, means a single or double-stranded polymer of deoxyribonucleotide or ribonucleotide bases and includes DNA and corresponding RNA molecules, including HnRNA and mRNA molecules, both sense and anti-sense strands, and comprehends cDNA, genomic DNA and recombinant DNA, as well as wholly or partially synthesized polynucleotides. An HnRNA molecule contains introns and corresponds to a DNA molecule in a generally one-to-one manner. An mRNA molecule corresponds to an HnRNA and DNA molecule from which the introns have been excised. A polynucleotide may consist of an entire gene, or any portion thereof. Operable anti-sense polynucleotides may comprise a fragment of the corresponding polynucleotide, and the definition of "polynucleotide" therefore includes all such operable anti-sense fragments.

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An "immunogenic portion" of an antigen is a portion that is capable of reacting with sera obtained from a Chlamydia-infected individual (i.e., generates an absorbance reading with sera from infected individuals that is at least three standard deviations above the absorbance obtained with sera from uninfected individuals, in a representative ELISA assay described herein). Such immunogenic portions generally comprise at least about 5 amino acid residues, more preferably at least about 10, and most preferably at least about 20 amino acid residues. Methods for preparing and identifying immunogenic portions of antigens of known sequence are well known in the art and include those summarized in Paul, Fundamental Immunology, 3rd ed., Raven Press, 1993, pp. 243-247 and references cited therein. Such techniques include screening polypeptides for the ability to react with antigen-specific antibodies, antisera and/or T-cell lines or clones. As used herein, antisera and antibodies are "antigenspecific" if they specifically bind to an antigen (i.e., they react with the protein in an ELISA or other immunoassay, and do not react detectably with unrelated proteins). Such antisera and antibodies may be prepared as described herein, and using well known techniques. An immunogenic portion of a native Chlamydia protein is a portion that reacts with such antisera and/or T-cells at a level that is not substantially less than the reactivity of the full length polypeptide (e.g., in an ELISA and/or T-cell reactivity assay). Such immunogenic portions may react within such assays at a level that is

similar to or greater than the reactivity of the full length polypeptide. Such screens may generally be performed using methods well known to those of ordinary skill in the art, such as those described in Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, 1988. For example, a polypeptide may be immobilized on a solid support and contacted with patient sera to allow binding of antibodies within the sera to the immobilized polypeptide. Unbound sera may then be removed and bound antibodies detected using, for example, ¹²⁵I-labeled Protein A.

Examples of immunogenic portions of antigens contemplated by the present invention include, for example, the T cell stimulating epitopes provided in SEQ ID NO: 9, 10, 18, 19, 31, 39, 93-96, 98, 100-102, 106, 108, 138-140, 158, 167, 168, 246, 247 and 254-256. Polypeptides comprising at least an immunogenic portion of one or more *Chlamydia* antigens as described herein may generally be used, alone or in combination, to detect Chlamydial infection in a patient.

The compositions and methods of the present invention also encompass variants of the above polypeptides and polynucleotide molecules. Such variants include, but are not limited to, naturally occurring allelic variants of the inventive sequences. In particular, variants include other *Chlamydiae* serovars, such as serovars D, E and F, as well as the several LGV serovars which share homology to the inventive polypeptide and polynucleotide molecules described herein. Preferably, the serovar homologues show 95-99% homology to the corresponding polypeptide sequence(s) described herein.

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A polypeptide "variant," as used herein, is a polypeptide that differs from the recited polypeptide only in conservative substitutions and/or modifications, such that the antigenic properties of the polypeptide are retained. In a preferred embodiment, variant polypeptides differ from an identified sequence by substitution, deletion or addition of five amino acids or fewer. Such variants may generally be identified by modifying one of the above polypeptide sequences, and evaluating the antigenic properties of the modified polypeptide using, for example, the representative procedures described herein. In other words, the ability of a variant to react with antigen-specific antisera may be enhanced or unchanged, relative to the native protein, or may be diminished by less than 50%, and preferably less than 20%, relative to the native protein. Such variants may generally be identified by modifying one of the above

51

polypeptide sequences and evaluating the reactivity of the modified polypeptide with antigen-specific antibodies or antisera as described herein. Preferred variants include those in which one or more portions, such as an N-terminal leader sequence or transmembrane domain, have been removed. Other preferred variants include variants in which a small portion (e.g., 1-30 amino acids, preferably 5-15 amino acids) has been removed from the N- and/or C-terminal of the mature protein.

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As used herein, a "conservative substitution" is one in which an amino acid is substituted for another amino acid that has similar properties, such that one skilled in the art of peptide chemistry would expect the secondary structure and hydropathic nature of the polypeptide to be substantially unchanged. Amino acid substitutions may generally be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity and/or the amphipathic nature of the residues. For example, negatively charged amino acids include aspartic acid and glutamic acid; positively charged amino acids include lysine and arginine; and amino acids with uncharged polar head groups having similar hydrophilicity values include leucine, isoleucine and valine; glycine and alanine; asparagine and glutamine; and serine, threonine, phenylalanine and tyrosine. Other groups of amino acids that may represent conservative changes include: (1) ala, pro, gly, glu, asp, gln, asn, ser, thr; (2) cys, ser, tyr, thr; (3) val, ile, leu, met, ala, phe; (4) lys, arg, his; and (5) phe, tyr, trp, his. A variant may also, or alternatively, contain nonconservative changes. In a preferred embodiment, variant polypeptides differ from a native sequence by substitution, deletion or addition of five amino acids or fewer. Variants may also (or alternatively) be modified by, for example, the deletion or addition of amino acids that have minimal influence on the immunogenicity, secondary structure and hydropathic nature of the polypeptide. Variants may also, or alternatively, contain other modifications, including the deletion or addition of amino acids that have minimal influence on the antigenic properties, secondary structure and hydropathic nature of the polypeptide. For example, a polypeptide may be conjugated to a signal (or leader) sequence at the N-terminal end of the protein which co-translationally or post-translationally directs transfer of the protein. The polypeptide may also be conjugated to a linker or other sequence for ease of synthesis, purification or identification of the polypeptide (e.g., poly-His), or to

enhance binding of the polypeptide to a solid support. For example, a polypeptide may be conjugated to an immunoglobulin Fc region.

52

A polynucleotide "variant" is a sequence that differs from the recited nucleotide sequence in having one or more nucleotide deletions, substitutions or additions such that the immunogenicity of the encoded polypeptide is not diminished, relative to the native protein. The effect on the immunogenicity of the encoded polypeptide may generally be assessed as described herein. Such modifications may be readily introduced using standard mutagenesis techniques, such as oligonucleotidedirected site-specific mutagenesis as taught, for example, by Adelman et al. (DNA, 2:183, 1983). Nucleotide variants may be naturally occurring allelic variants as discussed below, or non-naturally occurring variants. The polypeptides provided by the present invention include variants that are encoded by polynucleotide sequences which are substantially homologous to one or more of the polynucleotide sequences "Substantial homology," as used herein, refers to specifically recited herein. polynucleotide sequences that are capable of hybridizing under moderately stringent conditions. Suitable moderately stringent conditions include prewashing in a solution of 5X SSC, 0.5% SDS, 1.0 mM EDTA (pH 8.0); hybridizing at 50°C-65°C, 5X SSC, overnight or, in the event of cross-species homology, at 45°C with 0.5X SSC; followed by washing twice at 65°C for 20 minutes with each of 2X, 0.5X and 0.2X SSC containing 0.1% SDS. Such hybridizing polynucleotide sequences are also within the scope of this invention, as are nucleotide sequences that, due to code degeneracy, encode a polypeptide that is the same as a polypeptide of the present invention.

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Two nucleotide or polypeptide sequences are said to be "identical" if the sequence of nucleotides or amino acid residues in the two sequences is the same when aligned for maximum correspondence as described below. Comparisons between two sequences are typically performed by comparing the sequences over a comparison window to identify and compare local regions of sequence similarity. A "comparison window" as used herein, refers to a segment of at least about 20 contiguous positions, usually 30 to about 75, 40 to about 50, in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned.

WO 02/08267

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PCT/US01/23121

Optimal alignment of sequences for comparison may be conducted using the Megalign program in the Lasergene suite of bioinformatics software (DNASTAR, Inc., Madison, WI), using default parameters. This program embodies several alignment schemes described in the following references: Dayhoff, M.O. (1978) A model of evolutionary change in proteins - Matrices for detecting distant relationships. In Dayhoff, M.O. (ed.) Atlas of Protein Sequence and Structure, National Biomedical Resarch Foundaiton, Washington DC Vol. 5, Suppl. 3, pp. 345-358; Hein J. (1990) Unified Approach to Alignment and Phylogenes pp. 626-645 Methods in Enzymology vol. 183, Academic Press, Inc., San Diego, CA; Higgins, D.G. and Sharp, P.M. (1989) Fast and sensitive multiple sequence alignments on a microcomputer CABIOS 5:151-153; Myers, E.W. and Muller W. (1988) Optimal alignments in linear space CABIOS 4:11-17; Robinson, E.D. (1971) Comb. Theor 11:105; Santou, N. Nes, M. (1987) The neighbor joining method. A new method for reconstructing phylogenetic trees Mol. Biol. Evol. 4:406-425; Sneath, P.H.A. and Sokal, R.R. (1973) Numerical Taxonomy the Principles and Practice of Numerical Taxonomy, Freeman Press, San Francisco, CA; Wilbur, W.J. and Lipman, D.J. (1983) Rapid similarity searches of nucleic acid and protein data banks Proc. Natl. Acad., Sci. USA 80:726-730.

Alternatively, optimal alignment of sequences for comparison may be conducted by the local identity algorithm of Smith and Waterman (1981) *Add. APL. Math* 2:482, by the identity alignment algorithm of Needleman and Wunsch (1970) J. Mol. Biol. 48:443, by the search for similarity methods of Pearson and Lipman (1988) Proc. Natl. Acad. Sci. (U.S.A.) 85: 2444, by computerized implementations of these algorithms (GAP, BESTFIT, BLAST, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group (GCG), 575 Science Dr., Madison, WI), or by inspection.

One illustrative example of algorithms that are suitable for determining percent sequence identity and sequence similarity are the BLAST and BLAST 2.0 algorithms, which are described in Altschul et al. (1977) Nuc. Acids Res. 25:3389-3402 and Altschul et al. (1990) J. Mol. Biol. 215:403-410, respectively. BLAST and BLAST 2.0 can be used, for example with the parameters described herein, to determine percent sequence identity for the polynucleotides and polypeptides of the invention. Software for performing BLAST analyses is publicly available through the National Center for

WO 02/08267

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Biotechnology Information (http://www.ncbi.nlm.nih.gov/) In one illustrative example, cumulative scores can be calculated using, for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always >0) and N (penalty score for mismatching residues; always <0). For amino acid sequences, a scoring matrix can be used to calculate the cumulative score. Extension of the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength (W) of 11, and expectation (E) of 10, and the BLOSUM62 scoring matrix (see Henikoff and Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915) alignments, (B) of 50, expectation (E) of 10, M=5, N=-4 and a comparison of both strands.

54

PCT/US01/23121

Preferably, the "percentage of sequence identity" is determined by comparing two optimally aligned sequences over a window of comparison of at least 20 positions, wherein the portion of the polynucleotide or amino acid sequence in the comparison window may comprise additions or deletions (i.e. gaps) of 20 percent or less, usually 5 to 15 percent, or 10 to 12 percent, as compared to the reference sequences (which does not comprise additions or deletions) for optimal alignment of the two sequences. The percentage is calculated by determining the number of positions at which the identical nucleic acid bases or amino acid residue occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the reference sequence (i.e. the window size) and multiplying the results by 100 to yield the percentage of sequence identity.

Therefore, the present invention provides polynucleotide and polypeptide sequences having substantial identity to the sequences disclosed herein, for example those comprising at least 50% or more sequence identity, preferably at least 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, or 99% or higher, sequence identity compared to a polynucleotide or polypeptide sequence of this invention using the methods described herein, (e.g., BLAST analysis using standard parameters, as described below). One skilled in this art will recognize that these values can be

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appropriately adjusted to determine corresponding identity of proteins encoded by two polynucleotide sequences by taking into account codon degeneracy, amino acid similarity, reading frame positioning and the like.

In additional embodiments, the present invention provides isolated polynucleotides or polypeptides comprising various lengths of contiguous stretches of sequence identical to or complementary to one or more of the sequences disclosed herein. For example, polynucleotides and polypeptides encompassed by this invention may comprise at least about 15, 20, 30, 40, 50, 75, 100, 150, 200, 300, 400, 500 or 1000 or more contiguous nucleotides of one or more of the disclosed sequences, as well as all intermediate lengths therebetween. It will be readily understood that "intermediate lengths", in this context, means any length between the quoted values, such as 16, 17, 18, 19, etc.; 21, 22, 23, etc.; 30, 31, 32, etc.; 50, 51, 52, 53, etc.; 100, 101, 102, 103, etc.; 150, 151, 152, 153, etc.; including all integers through the 200-500; 500-1,000, and the like.

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The polynucleotides of the present invention, or fragments thereof, regardless of the length of the coding sequence itself, may be combined with other DNA sequences, such as promoters, polyadenylation signals, additional restriction enzyme sites, multiple cloning sites, other coding segments, and the like, such that their overall length may vary considerably. It is therefore contemplated that a nucleic acid fragment of almost any length may be employed, with the total length preferably being limited by the ease of preparation and use in the intended recombinant DNA protocol. For example, illustrative DNA segments with total lengths of about 10,000, about 5000, about 3000, about 2,000, about 1,000, about 500, about 200, about 100, about 50 base pairs in length, and the like, (including all intermediate lengths) are contemplated to be useful in many implementations of this invention.

Also included in the scope of the present invention are alleles of the genes encoding the nucleotide sequences recited in herein. As used herein, an "allele" or "allellic sequence" is an alternative form of the gene which may result from at least one mutation in the nucleic acid sequence. Alleles may result in altered mRNAs or polypeptides whose structure or function may or may not be altered. Any given gene may have none, one, or many allelic forms. Common mutational changes which give rise to alleles are generally ascribed to natural deletions, additions, or substitutions of

WO 02/08267

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nucleotides. Each of these types of changes may occur alone or in combination with the others, one or more times in a given sequence.

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In specific embodiments, the subject invention discloses polypeptides comprising at least an immunogenic portion of a Chlamydia antigen (or a variant of such an antigen), that comprises one or more of the amino acid sequences encoded by (a) a polynucleotide sequence selected from the group consisting of SEQ ID NO: 358-361, 407-430, 525-559, 582-598; (b) the complements of such DNA sequences or (c) DNA sequences substantially homologous to a sequence in (a) or (b). As discussed in the Examples below, several of the Chlamydia antigens disclosed herein recognize a T cell line that recognizes both Chlamydia trachomatis and Chlamydia pneumoniae infected monocyte-derived dendritic cells, indicating that they may represent an immunoreactive epitope shared by Chlamydia trachomatis and Chlamydia pneumoniae. The antigens may thus be employed in a vaccine for both C. trachomatis genital tract infections and for C. pneumonia infections. Further characterization of these Chlamydia antigens from Chlamydia trachomatis and Chlamydia pneumonia to determine the extent of cross-reactivity is provided in Example 6. Additionally, Example 4 describes cDNA fragments (SEQ ID NO: 15, 16 and 33) isolated from C. trachomatis which encode proteins (SEQ ID NO: 17-19 and 32) capable of stimulating a Chlamydiaspecific murine CD8+ T cell line.

In general, *Chlamydia* antigens, and polynucleotide sequences encoding such antigens, may be prepared using any of a variety of procedures. For example, polynucleotide molecules encoding *Chlamydia* antigens may be isolated from a *Chlamydia* genomic or cDNA expression library by screening with a *Chlamydia*-specific T cell line as described below, and sequenced using techniques well known to those of skill in the art. Additionally, a polynucleotide may be identified, as described in more detail below, by screening a microarray of cDNAs for *Chlamydia*-associated expression (*i.e.*, expression that is at least two fold greater in *Chlamydia*-infected cells than in controls, as determined using a representative assay provided herein). Such screens may be performed using a Synteni microarray (Palo Alto, CA) according to the manufacturer's instructions (and essentially as described by Schena et al., *Proc. Natl. Acad. Sci. USA 93*:10614-10619, 1996 and Heller et al., *Proc. Natl. Acad. Sci. USA 94*:2150-2155, 1997). Alternatively, polypeptides may be amplified from cDNA

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prepared from cells expressing the proteins described herein. Such polynucleotides may be amplified via polymerase chain reaction (PCR). For this approach, sequence-specific primers may be designed based on the sequences provided herein, and may be purchased or synthesized.

Antigens may be produced recombinantly, as described below, by inserting a polynucleotide sequence that encodes the antigen into an expression vector and expressing the antigen in an appropriate host. Antigens may be evaluated for a desired property, such as the ability to react with sera obtained from a *Chlamydia*-infected individual as described herein, and may be sequenced using, for example, traditional Edman chemistry. *See* Edman and Berg, *Eur. J. Biochem.* 80:116-132, 1967.

Polynucleotide sequences encoding antigens may also be obtained by screening an appropriate *Chlamydia* cDNA or genomic DNA library for polynucleotide sequences that hybridize to degenerate oligonucleotides derived from partial amino acid sequences of isolated antigens. Degenerate oligonucleotide sequences for use in such a screen may be designed and synthesized, and the screen may be performed, as described (for example) in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratories, Cold Spring Harbor, NY (and references cited therein). Polymerase chain reaction (PCR) may also be employed, using the above oligonucleotides in methods well known in the art, to isolate a nucleic acid probe from a cDNA or genomic library. The library screen may then be performed using the isolated probe.

An amplified portion may be used to isolate a full length gene from a suitable library (e.g., a Chlamydia cDNA library) using well known techniques. Within such techniques, a library (cDNA or genomic) is screened using one or more polynucleotide probes or primers suitable for amplification. Preferably, a library is size-selected to include larger molecules. Random primed libraries may also be preferred for identifying 5' and upstream regions of genes. Genomic libraries are preferred for obtaining introns and extending 5' sequences.

For hybridization techniques, a partial sequence may be labeled (e.g., by nick-translation or end-labeling with ³²P) using well known techniques. A bacterial or bacteriophage library is then screened by hybridizing filters containing denatured bacterial colonies (or lawns containing phage plaques) with the labeled probe (see

58

Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratories, Cold Spring Harbor, NY, 1989). Hybridizing colonies or plaques are selected and expanded, and the DNA is isolated for further analysis. cDNA clones may be analyzed to determine the amount of additional sequence by, for example, PCR using a primer from the partial sequence and a primer from the vector. Restriction maps and partial sequences may be generated to identify one or more overlapping clones. The complete sequence may then be determined using standard techniques, which may involve generating a series of deletion clones. The resulting overlapping sequences are then assembled into a single contiguous sequence. A full length cDNA molecule can be generated by ligating suitable fragments, using well known techniques.

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Alternatively, there are numerous amplification techniques for obtaining a full length coding sequence from a partial cDNA sequence. Within such techniques, amplification is generally performed via PCR. Any of a variety of commercially available kits may be used to perform the amplification step. Primers may be designed using techniques well known in the art (see, for example, Mullis et al., Cold Spring Harbor Symp. Quant. Biol. 51:263, 1987; Erlich ed., PCR Technology, Stockton Press, NY, 1989), and software well known in the art may also be employed. Primers are preferably 22-30 nucleotides in length, have a GC content of at least 50% and anneal to the target sequence at temperatures of about 68°C to 72°C. The amplified region may be sequenced as described above, and overlapping sequences assembled into a contiguous sequence.

One such amplification technique is inverse PCR (see Triglia et al., Nucl. Acids Res. 16:8186, 1988), which uses restriction enzymes to generate a fragment in the known region of the gene. The fragment is then circularized by intramolecular ligation and used as a template for PCR with divergent primers derived from the known region. Within an alternative approach, sequences adjacent to a partial sequence may be retrieved by amplification with a primer to a linker sequence and a primer specific to a known region. The amplified sequences are typically subjected to a second round of amplification with the same linker primer and a second primer specific to the known region. A variation on this procedure, which employs two primers that initiate extension in opposite directions from the known sequence, is described in WO 96/38591. Additional techniques include capture PCR (Lagerstrom et al., PCR Methods

Applic. 1:111-19, 1991) and walking PCR (Parker et al., Nucl. Acids. Res. 19:3055-60, 1991). Transcription-Mediated Amplification, or TMA is another method that may be utilized for the amplification of DNA, rRNA, or mRNA, as described in Patent No. PCT/US91/03184. This autocatalytic and isothermic non-PCR based method utilizes two primers and two enzymes: RNA polymerase and reverse transcriptase. One primer contains a promoter sequence for RNA polymerase. In the first amplification, the promoter-primer hybridizes to the target rRNA at a defined site. Reverse transcriptase creates a DNA copy of the target rRNA by extension from the 3'end of the promoterprimer. The RNA in the resulting complex is degraded and a second primer binds to the DNA copy. A new strand of DNA is synthesized from the end of the primer by reverse transcriptase creating double stranded DNA. RNA polymerase recognizes the promoter sequence in the DNA template and initiates transcription. Each of the newly synthesized RNA amplicons re-enters the TMA process and serves as a template for a new round of replication leading to the expotential expansion of the RNA amplicon. Other methods employing amplification may also be employed to obtain a full length cDNA sequence.

In certain instances, it is possible to obtain a full length cDNA sequence by analysis of sequences provided in an expressed sequence tag (EST) database, such as that available from GenBank. Searches for overlapping ESTs may generally be performed using well known programs (e.g., NCBI BLAST searches), and such ESTs may be used to generate a contiguous full length sequence. Full length cDNA sequences may also be obtained by analysis of genomic fragments.

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Polynucleotide variants may generally be prepared by any method known in the art, including chemical synthesis by, for example, solid phase phosphoramidite chemical synthesis. Modifications in a polynucleotide sequence may also be introduced using standard mutagenesis techniques, such as oligonucleotide-directed site-specific mutagenesis (see Adelman et al., DNA 2:183, 1983). Alternatively, RNA molecules may be generated by in vitro or in vivo transcription of DNA sequences encoding a Chlamydial protein, or portion thereof, provided that the DNA is incorporated into a vector with a suitable RNA polymerase promoter (such as T7 or SP6). Certain portions may be used to prepare an encoded polypeptide, as described herein. In addition, or alternatively, a portion may be administered to a patient such that the encoded

polypeptide is generated *in vivo* (e.g., by transfecting antigen-presenting cells, such as dendritic cells, with a cDNA construct encoding a *Chlamydial* polypeptide, and administering the transfected cells to the patient).

A portion of a sequence complementary to a coding sequence (i.e., an antisense polynucleotide) may also be used as a probe or to modulate gene expression. cDNA constructs that can be transcribed into antisense RNA may also be introduced into cells of tissues to facilitate the production of antisense RNA. An antisense polynucleotide may be used, as described herein, to inhibit expression of a *Chlamydial* protein. Antisense technology can be used to control gene expression through triplehelix formation, which compromises the ability of the double helix to open sufficiently for the binding of polymerases, transcription factors or regulatory molecules (see Gee et al., In Huber and Carr, Molecular and Immunologic Approaches, Futura Publishing Co. (Mt. Kisco, NY; 1994)). Alternatively, an antisense molecule may be designed to hybridize with a control region of a gene (e.g., promoter, enhancer or transcription initiation site), and block transcription of the gene; or to block translation by inhibiting binding of a transcript to ribosomes.

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A portion of a coding sequence, or of a complementary sequence, may also be designed as a probe or primer to detect gene expression. Probes may be labeled with a variety of reporter groups, such as radionuclides and enzymes, and are preferably at least 10 nucleotides in length, more preferably at least 20 nucleotides in length and still more preferably at least 30 nucleotides in length. Primers, as noted above, are preferably 22-30 nucleotides in length.

Any polynucleotide may be further modified to increase stability *in vivo*. Possible modifications include, but are not limited to, the addition of flanking sequences at the 5' and/or 3' ends; the use of phosphorothioate or 2' O-methyl rather than phosphodiesterase linkages in the backbone; and/or the inclusion of nontraditional bases such as inosine, queosine and wybutosine, as well as acetyl- methyl-, thio- and other modified forms of adenine, cytidine, guanine, thymine and uridine.

Nucleotide sequences as described herein may be joined to a variety of other nucleotide sequences using established recombinant DNA techniques. For example, a polynucleotide may be cloned into any of a variety of cloning vectors, including plasmids, phagemids, lambda phage derivatives and cosmids. Vectors of

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particular interest include expression vectors, replication vectors, probe generation vectors and sequencing vectors. In general, a vector will contain an origin of replication functional in at least one organism, convenient restriction endonuclease sites and one or more selectable markers. Other elements will depend upon the desired use, and will be apparent to those of ordinary skill in the art.

Synthetic polypeptides having fewer than about 100 amino acids, and generally fewer than about 50 amino acids, may be generated using techniques well known in the art. For example, such polypeptides may be synthesized using any of the commercially available solid-phase techniques, such as the Merrifield solid-phase synthesis method, where amino acids are sequentially added to a growing amino acid chain. See Merrifield, J. Am. Chem. Soc. 85:2149-2146, 1963. Equipment for automated synthesis of polypeptides is commercially available from suppliers such as Perkin Elmer/Applied BioSystems Division, Foster City, CA, and may be operated according to the manufacturer's instructions.

As noted above, immunogenic portions of *Chlamydia* antigens may be prepared and identified using well known techniques, such as those summarized in Paul, *Fundamental Immunology*, 3d ed., Raven Press, 1993, pp. 243-247 and references cited therein. Such techniques include screening polypeptide portions of the native antigen for immunogenic properties. The representative ELISAs described herein may generally be employed in these screens. An immunogenic portion of a polypeptide is a portion that, within such representative assays, generates a signal in such assays that is substantially similar to that generated by the full length antigen. In other words, an immunogenic portion of a *Chlamydia* antigen generates at least about 20%, and preferably about 100%, of the signal induced by the full length antigen in a model ELISA as described herein.

Portions and other variants of *Chlamydia* antigens may be generated by synthetic or recombinant means. Variants of a native antigen may generally be prepared using standard mutagenesis techniques, such as oligonucleotide-directed site-specific mutagenesis. Sections of the polynucleotide sequence may also be removed using standard techniques to permit preparation of truncated polypeptides.

Recombinant polypeptides containing portions and/or variants of a native antigen may be readily prepared from a polynucleotide sequence encoding the

62

polypeptide using a variety of techniques well known to those of ordinary skill in the art. For example, supernatants from suitable host/vector systems which secrete recombinant protein into culture media may be first concentrated using a commercially available filter. Following concentration, the concentrate may be applied to a suitable purification matrix such as an affinity matrix or an ion exchange resin. Finally, one or more reverse phase HPLC steps can be employed to further purify a recombinant protein.

Any of a variety of expression vectors known to those of ordinary skill in the art may be employed to express recombinant polypeptides as described herein. Expression may be achieved in any appropriate host cell that has been transformed or transfected with an expression vector containing a polynucleotide molecule that encodes a recombinant polypeptide. Suitable host cells include prokaryotes, yeast and higher eukaryotic cells. Preferably, the host cells employed are *E. coli*, yeast or a mammalian cell line, such as COS or CHO. The DNA sequences expressed in this manner may encode naturally occurring antigens, portions of naturally occurring antigens, or other variants thereof.

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In general, regardless of the method of preparation, the polypeptides disclosed herein are prepared in an isolated, substantially pure, form. Preferably, the polypeptides are at least about 80% pure, more preferably at least about 90% pure and most preferably at least about 99% pure.

Within certain specific embodiments, a polypeptide may be a fusion protein that comprises multiple polypeptides as described herein, or that comprises at least one polypeptide as described herein and an unrelated sequence, such as a known *Chlamydial* protein. A fusion partner may, for example, assist in providing T helper epitopes (an immunological fusion partner), preferably T helper epitopes recognized by humans, or may assist in expressing the protein (an expression enhancer) at higher yields than the native recombinant protein. Certain preferred fusion partners are both immunological and expression enhancing fusion partners. Other fusion partners may be selected so as to increase the solubility of the protein or to enable the protein to be targeted to desired intracellular compartments. Still further fusion partners include affinity tags, which facilitate purification of the protein. A DNA sequence encoding a fusion protein of the present invention may be constructed using known recombinant

DNA techniques to assemble separate DNA sequences encoding, for example, the first and second polypeptides, into an appropriate expression vector. The 3' end of a DNA sequence encoding the first polypeptide is ligated, with or without a peptide linker, to the 5' end of a DNA sequence encoding the second polypeptide so that the reading frames of the sequences are in phase to permit mRNA translation of the two DNA sequences into a single fusion protein that retains the biological activity of both the first and the second polypeptides.

A peptide linker sequence may be employed to separate the first and the second polypeptides by a distance sufficient to ensure that each polypeptide folds into its secondary and tertiary structures. Such a peptide linker sequence is incorporated into the fusion protein using standard techniques well known in the art. Suitable peptide linker sequences may be chosen based on the following factors: (1) their ability to adopt a flexible extended conformation; (2) their inability to adopt a secondary structure that could interact with functional epitopes on the first and second polypeptides; and (3) the lack of hydrophobic or charged residues that might react with the polypeptide functional epitopes. Preferred peptide linker sequences contain Gly, Asn and Ser residues. Other near neutral amino acids, such as Thr and Ala may also be used in the linker sequence. Amino acid sequences which may be usefully employed as linkers include those disclosed in Maratea et al., Gene 40:39-46, 1985; Murphy et al., Proc. Natl. Acad. Sci. USA 83:8258-8562, 1986; U.S. Patent No. 4,935,233 and U.S. Patent No. 4,751,180. The linker sequence may be from 1 to about 50 amino acids in length. As an alternative to the use of a peptide linker sequence (when desired), one can utilize non-essential N-terminal amino acid regions (when present) on the first and second polypeptides to separate the functional domains and prevent steric hindrance.

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The ligated DNA sequences are operably linked to suitable transcriptional or translational regulatory elements. The regulatory elements responsible for expression of DNA are located only 5' to the DNA sequence encoding the first polypeptides. Similarly, stop codons required to end translation and transcription termination signals are only present 3' to the DNA sequence encoding the second polypeptide.

Fusion proteins are also provided that comprise a polypeptide of the present invention together with an unrelated immunogenic protein. Preferably the

immunogenic protein is capable of eliciting a recall response. Examples of such proteins include tetanus, tuberculosis and hepatitis proteins (see, for example, Stoute et al. New Engl. J. Med., 336:86-91, 1997).

Within preferred embodiments, an immunological fusion partner is derived from protein D, a surface protein of the gram-negative bacterium Haemophilus influenza B (WO 91/18926). Preferably, a protein D derivative comprises approximately the first third of the protein (e.g., the first N-terminal 100-110 amino acids), and a protein D derivative may be lipidated. Within certain preferred embodiments, the first 109 residues of a Lipoprotein D fusion partner is included on the N-terminus to provide the polypeptide with additional exogenous T-cell epitopes and to increase the expression level in E. coli (thus functioning as an expression enhancer). The lipid tail ensures optimal presentation of the antigen to antigen presenting cells. Other fusion partners include the non-structural protein from influenzae virus, NS1 (hemaglutinin). Typically, the N-terminal 81 amino acids are used, although different fragments that include T-helper epitopes may be used.

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In another embodiment, the immunological fusion partner is the protein known as LYTA, or a portion thereof (preferably a C-terminal portion). LYTA is derived from *Streptococcus pneumoniae*, which synthesizes an N-acetyl-L-alanine amidase known as amidase LYTA (encoded by the LytA gene; *Gene 43*:265-292, 1986). LYTA is an autolysin that specifically degrades certain bonds in the peptidoglycan backbone. The C-terminal domain of the LYTA protein is responsible for the affinity to the choline or to some choline analogues such as DEAE. This property has been exploited for the development of *E. coli* C-LYTA expressing plasmids useful for expression of fusion proteins. Purification of hybrid proteins containing the C-LYTA fragment at the amino terminus has been described (*see Biotechnology 10*:795-798, 1992). Within a preferred embodiment, a repeat portion of LYTA may be incorporated into a fusion protein. A repeat portion is found in the C-terminal region starting at residue 178. A particularly preferred repeat portion incorporates residues 188-305.

In another embodiment, a Mycobacterium tuberculosis-derived Ra12 polynucleotide is linked to at least an immunogenic portion of a polynucleotide of this invention. Ra12 compositions and methods for their use inenhancing expression of heterologous polynucleotide sequences is described in U.S. Patent Application

60/158,585, the disclosure of which is incorporated herein by reference in its entirety. Briefly, Ra12 refers to a polynucleotide region that is a subsequence of a *Mycobacterium tuberculosis* MTB32A nucleic acid. MTB32A is a serine protease of 32 KD molecular weight encoded by a gene in virulent and avirulent strains of *M. tuberculosis*. The nucleotide sequence and amino acid sequence of MTB32A have been described (U.S. Patent Application 60/158,585; see also, Skeiky *et al.*, *Infection and Immun.* (1999) 67:3998-4007, incorporated herein by reference. In one embodiment, the Ra12 polypeptide used in the production of fusion polypeptides comprises a C-terminal fragment of the MTB32A coding sequence that is effective for enhancing the expression and/or immunogenicity of heterologous Chlamydial antigenic polypeptides with which it is fused. In another embodiment, the Ra12 polypeptide corresponds to an approximately 14 kD. C-terminal fragment of MTB32A comprising some or all of amino acid residues 192 to 323 of MTB32A.

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Recombinant nucleic acids, which encode a fusion polypeptide comprising a Ra12 polypeptide and a heterologous Chlamydia polypeptide of interest, can be readily constructed by conventional genetic engineering techniques. Recombinant nucleic acids are constructed so that, preferably, a Ra12 polynucleotide sequence is located 5' to a selected heterologous Chlamydia polynucleotide sequence. It may also be appropriate to place a Ra12 polynucleotide sequence 3' to a selected heterologous polynucleotide sequence or to insert a heterologous polynucleotide sequence into a site within a Ra12 polynucleotide sequence.

In addition, any suitable polynucleotide that encodes a Ra12 or a portion or other variant thereof can be used in constructing recombinant fusion polynucleotides comprising Ra12 and one or more Chlamydia polynucleotides disclosed herein. Preferred Ra12 polynucleotides generally comprise at least about 15 consecutive nucleotides, at least about 30 nucleotides, at least about 60 nucleotides, at least about 100 nucleotides, at least about 200 nucleotides, or at least about 300 nucleotides that encode a portion of a Ra12 polypeptide.

Ra12 polynucleotides may comprise a native sequence (i.e., an endogenous sequence that encodes a Ra12 polypeptide or a portion thereof) or may comprise a variant of such a sequence. Ra12 polynucleotide variants may contain one

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or more substitutions, additions, deletions and/or insertions such that the biological activity of the encoded fusion polypeptide is not substantially diminished, relative to a fusion polypeptide comprising a native Ra12 polypeptide. Variants preferably exhibit at least about 70% identity, more preferably at least about 80% identity and most preferably at least about 90% identity to a polynucleotide sequence that encodes a native Ra12 polypeptide or a portion thereof.

In another aspect, the present invention provides methods for using one or more of the above polypeptides or fusion proteins (or polynucleotides encoding such polypeptides or fusion proteins) to induce protective immunity against Chlamydial infection in a patient. As used herein, a "patient" refers to any warm-blooded animal, preferably a human. A patient may be afflicted with a disease, or may be free of detectable disease and/or infection. In other words, protective immunity may be induced to prevent or treat Chlamydial infection.

In this aspect, the polypeptide, fusion protein or polynucleotide molecule is generally present within a pharmaceutical composition or a vaccine. Pharmaceutical compositions may comprise one or more polypeptides, each of which may contain one or more of the above sequences (or variants thereof), and a physiologically acceptable carrier. Vaccines may comprise one or more of the above polypeptides and an immunostimulant, such as an adjuvant or a liposome (into which the polypeptide is incorporated). Such pharmaceutical compositions and vaccines may also contain other *Chlamydia* antigens, either incorporated into a combination polypeptide or present within a separate polypeptide.

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Alternatively, a vaccine may contain polynucleotides encoding one or more polypeptides or fusion proteins as described above, such that the polypeptide is generated *in situ*. In such vaccines, the polynucleotides may be present within any of a variety of delivery systems known to those of ordinary skill in the art, including nucleic acid expression systems, bacterial and viral expression systems. Appropriate nucleic acid expression systems contain the necessary polynucleotide sequences for expression in the patient (such as a suitable promoter and terminating signal). Bacterial delivery systems involve the administration of a bacterium (such as *Bacillus-Calmette-Guerrin*) that expresses an immunogenic portion of the polypeptide on its cell surface. In a

67

preferred embodiment, the polynucleotides may be introduced using a viral expression system (e.g., vaccinia or other pox virus, retrovirus, or adenovirus), which may involve the use of a non-pathogenic (defective) virus. Techniques for incorporating polynucleotides into such expression systems are well known to those of ordinary skill in the art. The polynucleotides may also be administered as "naked" plasmid vectors as described, for example, in Ulmer et al., Science 259:1745-1749, 1993 and reviewed by Cohen, Science 259:1691-1692, 1993. Techniques for incorporating DNA into such vectors are well known to those of ordinary skill in the art. A retroviral vector may additionally transfer or incorporate a gene for a selectable marker (to aid in the identification or selection of transduced cells) and/or a targeting moiety, such as a gene that encodes a ligand for a receptor on a specific target cell, to render the vector target specific. Targeting may also be accomplished using an antibody, by methods known to those of ordinary skill in the art.

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Other formulations for therapeutic purposes include colloidal dispersion systems, such as macromolecule complexes, nanocapsules, microspheres, beads, and lipid-based systems including oil-in-water emulsions, micelles, mixed micelles, and liposomes. A preferred colloidal system for use as a delivery vehicle *in vitro* and *in vivo* is a liposome (*i.e.*, an artificial membrane vesicle). The uptake of naked polynucleotides may be increased by incorporating the polynucleotides into and/or onto biodegradable beads, which are efficiently transported into the cells. The preparation and use of such systems is well known in the art.

In a related aspect, a polynucleotide vaccine as described above may be administered simultaneously with or sequentially to either a polypeptide of the present invention or a known *Chlamydia* antigen. For example, administration of polynucleotides encoding a polypeptide of the present invention, either "naked" or in a delivery system as described above, may be followed by administration of an antigen in order to enhance the protective immune effect of the vaccine.

Polypeptides and polynucleotides disclosed herein may also be employed in adoptive immunotherapy for the treatment of *Chlamydial* infection. Adoptive immunotherapy may be broadly classified into either active or passive immunotherapy. In active immunotherapy, treatment relies on the *in vivo* stimulation of the endogenous

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host immune system with the administration of immune response-modifying agents (for example, vaccines, bacterial adjuvants, and/or cytokines).

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In passive immunotherapy, treatment involves the delivery of biologic reagents with established immune reactivity (such as effector cells or antibodies) that can directly or indirectly mediate anti-Chlamydia effects and does not necessarily depend on an intact host immune system. Examples of effector cells include T lymphocytes (for example, CD8+ cytotoxic T-lymphocyte, CD4+ T-helper), killer cells (such as Natural Killer cells, lymphokine-activated killer cells), B cells, or antigen presenting cells (such as dendritic cells and macrophages) expressing the disclosed antigens. The polypeptides disclosed herein may also be used to generate antibodies or anti-idiotypic antibodies (as in U.S. Patent No. 4,918,164), for passive immunotherapy.

The predominant method of procuring adequate numbers of T-cells for adoptive immunotherapy is to grow immune T-cells in vitro. Culture conditions for expanding single antigen-specific T-cells to several billion in number with retention of antigen recognition in vivo are well known in the art. These in vitro culture conditions typically utilize intermittent stimulation with antigen, often in the presence of cytokines, such as IL-2, and non-dividing feeder cells. As noted above, the immunoreactive polypeptides described herein may be used to rapidly expand antigen-specific T cell cultures in order to generate sufficient number of cells for immunotherapy. particular, antigen-presenting cells, such as dendritic, macrophage, monocyte, fibroblast, or B-cells, may be pulsed with immunoreactive polypeptides, or polynucleotide sequence(s) may be introduced into antigen presenting cells, using a variety of standard techniques well known in the art. For example, antigen presenting cells may be transfected or transduced with a polynucleotide sequence, wherein said sequence contains a promoter region appropriate for increasing expression, and can be expressed as part of a recombinant virus or other expression system. Several viral vectors may be used to transduce an antigen presenting cell, including pox virus, vaccinia virus, and adenovirus; also, antigen presenting cells may be transfected with polynucleotide sequences disclosed herein by a variety of means, including gene-gun technology, lipid-mediated delivery, electroporation, osmotic shock, and particlate delivery mechanisms, resulting in efficient and acceptable expression levels as determined by one of ordinary skill in the art. For cultured T-cells to be effective in

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therapy, the cultured T-cells must be able to grow and distribute widely and to survive long term *in vivo*. Studies have demonstrated that cultured T-cells can be induced to grow *in vivo* and to survive long term in substantial numbers by repeated stimulation with antigen supplemented with IL-2 (see, for example, Cheever, M., *et al*, "Therapy With Cultured T Cells: Principles Revisited," *Immunological Reviews*, 157:177, 1997).

The polypeptides disclosed herein may also be employed to generate and/or isolate chlamydial-reactive T-cells, which can then be administered to the patient. In one technique, antigen-specific T-cell lines may be generated by *in vivo* immunization with short peptides corresponding to immunogenic portions of the disclosed polypeptides. The resulting antigen specific CD8+ or CD4+ T-cell clones may be isolated from the patient, expanded using standard tissue culture techniques, and returned to the patient.

Alternatively, peptides corresponding to immunogenic portions of the polypeptides may be employed to generate *Chlamydia* reactive T cell subsets by selective *in vitro* stimulation and expansion of autologous T cells to provide antigenspecific T cells which may be subsequently transferred to the patient as described, for example, by Chang *et al*, (*Crit. Rev. Oncol. Hematol.*, 22(3), 213, 1996). Cells of the immune system, such as T cells, may be isolated from the peripheral blood of a patient, using a commercially available cell separation system, such as IsolexTM System, available from Nexell Therapeutics, Inc. Irvine, CA. The separated cells are stimulated with one or more of the immunoreactive polypeptides contained within a delivery vehicle, such as a microsphere, to provide antigen-specific T cells. The population of antigen-specific T cells is then expanded using standard techniques and the cells are administered back to the patient.

In other embodiments, T-cell and/or antibody receptors specific for the polypeptides disclosed herein can be cloned, expanded, and transferred into other vectors or effector cells for use in adoptive immunotherapy. In particular, T cells may be transfected with the appropriate genes to express the variable domains from chlamydia specific monoclonal antibodies as the extracellular recognition elements and joined to the T cell receptor signaling chains, resulting in T cell activation, specific lysis, and cytokine release. This enables the T cell to redirect its specificity in an MHC-independent manner. See for example, Eshhar, Z., Cancer Immunol Immunother, 45(3-

4):131-6, 1997 and Hwu, P., et al, *Cancer Res*, 55(15):3369-73, 1995. Another embodiment may include the transfection of chlamydia antigen specific alpha and beta T cell receptor chains into alternate T cells, as in Cole, DJ, et al, *Cancer Res*, 55(4):748-52, 1995.

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In a further embodiment, syngeneic or autologous dendritic cells may be pulsed with peptides corresponding to at least an immunogenic portion of a polypeptide disclosed herein. The resulting antigen-specific dendritic cells may either be transferred into a patient, or employed to stimulate T cells to provide antigen-specific T cells which may, in turn, be administered to a patient. The use of peptide-pulsed dendritic cells to generate antigen-specific T cells and the subsequent use of such antigen-specific T cells to eradicate disease in a murine model has been demonstrated by Cheever et al, *Immunological Reviews*, 157:177, 1997). Additionally, vectors expressing the disclosed polynucleotides may be introduced into stem cells taken from the patient and clonally propagated *in vitro* for autologous transplant back into the same patient.

Within certain aspects, polypeptides, polynucleotides, T cells and/or binding agents disclosed herein may be incorporated into pharmaceutical compositions or immunogenic compositions (i.e., vaccines). Alternatively, a pharmaceutical composition may comprise an antigen-presenting cell (e.g. a dendritic cell) transfected with a Chlamydial polynucleotide such that the antigen presenting cell expresses a Chlamydial polypeptide. Pharmaceutical compositions comprise one or more such compounds and a physiologically acceptable carrier. Vaccines may comprise one or more such compounds and an immunostimulant. An immunostimulant may be any substance that enhances or potentiates an immune response to an exogenous antigen. Examples of immunostimulants include adjuvants, biodegradable microspheres (e.g., polylactic galactide) and liposomes (into which the compound is incorporated; see e.g., Fullerton, U.S. Patent No. 4,235,877). Vaccine preparation is generally described in, for example, M.F. Powell and M.J. Newman, eds., "Vaccine Design (the subunit and adjuvant approach)," Plenum Press (NY, 1995). Pharmaceutical compositions and vaccines within the scope of the present invention may also contain other compounds, which may be biologically active or inactive. For example, one or more immunogenic portions of other Chlamydial antigens may be present, either incorporated into a fusion polypeptide or as a separate compound, within the composition or vaccine.

A pharmaceutical composition or vaccine may contain DNA encoding one or more of the polypeptides as described above, such that the polypeptide is generated in situ. As noted above, the DNA may be present within any of a variety of delivery systems known to those of ordinary skill in the art, including nucleic acid expression systems, bacteria and viral expression systems. Numerous gene delivery techniques are well known in the art, such as those described by Rolland, Crit. Rev. Therap. Drug Carrier Systems 15:143-198, 1998, and references cited therein. Appropriate nucleic acid expression systems contain the necessary DNA sequences for expression in the patient (such as a suitable promoter and terminating signal). Bacterial delivery systems involve the administration of a bacterium (such as Bacillus-Calmette-Guerrin) that expresses an immunogenic portion of the polypeptide on its cell surface or secretes such an epitope.

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In a preferred embodiment, the DNA may be introduced using a viral expression system (e.g., vaccinia or other pox virus, retrovirus, adenovirus, baculovirus, togavirus, bacteriophage, and the like), which often involves the use of a non-pathogenic (defective), replication competent virus.

For example, many viral expression vectors are derived from viruses of the retroviridae family. This family includes the murine leukemia viruses, the mouse mammary tumor viruses, the human foamy viruses, Rous sarcoma virus, and the immunodeficiency viruses, including human, simian, and feline. Considerations when designing retroviral expression vectors are discussed in Comstock *et al.* (1997).

Excellent murine leukemia virus (MLV)-based viral expression vectors have been developed by Kim et al. (1998). In creating the MLV vectors, Kim et al. found that the entire gag sequence, together with the immediate upstream region, could be deleted without significantly affecting viral packaging or gene expression. Further, it was found that nearly the entire U3 region could be replaced with the immediately-early promoter of human cytomegalovirus without deleterious effects. Additionally, MCR and internal ribosome entry sites (IRES) could be added without adverse effects. Based on their observations, Kim et al. have designed a series of MLV-based expression vectors comprising one or more of the features described above.

As more has been learned about human foamy virus (HFV), characteristics of HFV that are favorable for its use as an expression vector have been

72

discovered. These characteristics include the expression of pol by splicing and start of translation at a defined initiation codon. Other aspects of HFV viral expression vectors are reviewed in Bodem *et al.* (1997).

Murakami et al. (1997) describe a Rous sarcoma virus (RSV)-based replication-competent avian retrovirus vectors, IR1 and IR2 to express a heterologous gene at a high level. In these vectors, the IRES derived from encephalomyocarditis virus (EMCV) was inserted between the env gene and the heterologous gene. The IR1 vector retains the splice-acceptor site that is present downstream of the env gene while the IR2 vector lacks it. Murakami et al. have shown high level expression of several different heterologous genes by these vectors.

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Recently, a number of lentivirus-based retroviral expression vectors have been developed. Kafri *et al.* (1997) have shown sustained expression of genes delivered directly into liver and muscle by a human immunodeficiency virus (HIV)-based expression vector. One benefit of the system is the inherent ability of HIV to transduce non-dividing cells. Because the viruses of Kafri *et al.* are pseudotyped with vesicular stomatitis virus G glycoprotein (VSVG), they can transduce a broad range of tissues and cell types.

A large number of adenovirus-based expression vectors have been developed, primarily due to the advantages offered by these vectors in gene therapy applications. Adenovirus expression vectors and methods of using such vectors are the subject of a number of United States patents, including United States Patent No. 5,698,202, United States Patent No. 5,616,326, United States Patent No. 5,585,362, and United States Patent No. 5,518,913, all incorporated herein by reference.

Additional adenoviral constructs are described in Khatri et al. (1997) and Tomanin et al. (1997). Khatri et al. describe novel ovine adenovirus expression vectors and their ability to infect bovine nasal turbinate and rabbit kidney cells as well as a range of human cell type, including lung and foreskin fibroblasts as well as liver, prostate, breast, colon and retinal lines. Tomanin et al. describe adenoviral expression vectors containing the T7 RNA polymerase gene. When introduced into cells containing a heterologous gene operably linked to a T7 promoter, the vectors were able to drive gene expression from the T7 promoter. The authors suggest that this system may be useful for the cloning and expression of genes encoding cytotoxic proteins.

Poxviruses are widely used for the expression of heterologous genes in mammalian cells. Over the years, the vectors have been improved to allow high expression of the heterologous gene and simplify the integration of multiple heterologous genes into a single molecule. In an effort to diminish cytopathic effects and to increase safety, vaccinia virus mutant and other poxviruses that undergo abortive infection in mammalian cells are receiving special attention (Oertli et al., 1997). The use of poxviruses as expression vectors is reviewed in Carroll and Moss (1997).

Togaviral expression vectors, which includes alphaviral expression vectors have been used to study the structure and function of proteins and for protein production purposes. Attractive features of togaviral expression vectors are rapid and efficient gene expression, wide host range, and RNA genomes (Huang, 1996). Also, recombinant vaccines based on alphaviral expression vectors have been shown to induce a strong humoral and cellular immune response with good immunological memory and protective effects (Tubulekas *et al.*, 1997). Alphaviral expression vectors and their use are discussed, for example, in Lundstrom (1997).

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In one study, Li and Garoff (1996) used Semliki Forest virus (SFV) expression vectors to express retroviral genes and to produce retroviral particles in BHK-21 cells. The particles produced by this method had protease and reverse transcriptase activity and were infectious. Furthermore, no helper virus could be detected in the virus stocks. Therefore, this system has features that are attractive for its use in gene therapy protocols.

Baculoviral expression vectors have traditionally been used to express heterologous proteins in insect cells. Examples of proteins include mammalian chemokine receptors (Wang et al., 1997), reporter proteins such as green fluorescent protein (Wu et al., 1997), and FLAG fusion proteins (Wu et al., 1997; Koh et al., 1997). Recent advances in baculoviral expression vector technology, including their use in virion display vectors and expression in mammalian cells is reviewed by Possee (1997). Other reviews on baculoviral expression vectors include Jones and Morikawa (1996) and O'Reilly (1997).

Other suitable viral expression systems are disclosed, for example, in Fisher-Hoch et al., *Proc. Natl. Acad. Sci. USA 86*:317-321, 1989; Flexner et al., *Ann. N.Y. Acad. Sci. 569*:86-103, 1989; Flexner et al., *Vaccine 8*:17-21, 1990; U.S. Patent

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PCT/US01/23121

Nos. 4,603,112, 4,769,330, and 5,017,487; WO 89/01973; U.S. Patent No. 4,777,127; GB 2,200,651; EP 0,345,242; WO 91/02805; Berkner, Biotechniques 6:616-627, 1988; Rosenfeld et al., Science 252:431-434, 1991; Kolls et al., Proc. Natl. Acad. Sci. USA 91:215-219, 1994; Kass-Eisler et al., Proc. Natl. Acad. Sci. USA 90:11498-11502, 1993; Guzman et al., Circulation 88:2838-2848, 1993; and Guzman et al., Cir. Res. 73:1202-1207, 1993. Techniques for incorporating DNA into such expression systems are well known to those of ordinary skill in the art. In other systems, the DNA may be introduced as "naked" DNA, as described, for example, in Ulmer et al., Science 259:1745-1749, 1993 and reviewed by Cohen, Science 259:1691-1692, 1993. The uptake of naked DNA may be increased by coating the DNA onto biodegradable beads, which are efficiently transported into the cells.

It will be apparent that a vaccine may comprise a polynucleotide and/or a polypeptide component, as desired. It will also be apparent that a vaccine may contain pharmaceutically acceptable salts of the polynucleotides and/or polypeptides provided herein. Such salts may be prepared from pharmaceutically acceptable non-toxic bases, including organic bases (e.g., salts of primary, secondary and tertiary amines and basic amino acids) and inorganic bases (e.g., sodium, potassium, lithium, ammonium, calcium and magnesium salts). While any suitable carrier known to those of ordinary skill in the art may be employed in the pharmaceutical compositions of this invention, the type of carrier will vary depending on the mode of administration. Compositions of the present invention may be formulated for any appropriate manner of administration, including for example, topical, oral, nasal, intravenous, intracranial, intraperitoneal, subcutaneous or intramuscular administration. For parenteral administration, such as subcutaneous injection, the carrier preferably comprises water, saline, alcohol, a fat, a wax or a buffer. For oral administration, any of the above carriers or a solid carrier, such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, sucrose, and magnesium carbonate, may be employed. Biodegradable microspheres (e.g., polylactate polyglycolate) may also be employed as carriers for the pharmaceutical compositions of this invention. Suitable biodegradable microspheres are disclosed, for example, in U.S. Patent Nos. 4,897,268 and 5,075,109.

Such compositions may also comprise buffers (e.g., neutral buffered saline or phosphate buffered saline), carbohydrates (e.g., glucose, mannose, sucrose or

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WO 02/08267 PCT/US01/23121

dextrans), mannitol, proteins, polypeptides or amino acids such as glycine, antioxidants, bacteriostats, chelating agents such as EDTA or glutathione, adjuvants (e.g., aluminum hydroxide), solutes that render the formulation isotonic, hypotonic or weakly hypertonic with the blood of a recipient, suspending agents, thickening agents and/or preservatives. Alternatively, compositions of the present invention may be formulated as a lyophilizate. Compounds may also be encapsulated within liposomes using well known technology.

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Any of a variety of immunostimulants may be employed in the vaccines of this invention. For example, an adjuvant may be included. Most adjuvants contain a substance designed to protect the antigen from rapid catabolism, such as aluminum hydroxide or mineral oil, and a stimulator of immune responses, such as lipid A, Bortadella pertussis or Mycobacterium tuberculosis derived proteins. Suitable adjuvants are commercially available as, for example, Freund's Incomplete Adjuvant and Complete Adjuvant (Difco Laboratories, Detroit, MI); Merck Adjuvant 65 (Merck and Company, Inc., Rahway, NJ); AS-2 (SmithKline Beecham, Philadelphia, PA); aluminum salts such as aluminum hydroxide gel (alum) or aluminum phosphate; salts of calcium, iron or zinc; an insoluble suspension of acylated tyrosine; acylated sugars; cationically anionically derivatized polysaccharides; polyphosphazenes; or biodegradable microspheres; monophosphoryl lipid A and quil A. Cytokines, such as GM-CSF or interleukin-2, -7, or -12, may also be used as adjuvants.

Within the vaccines provided herein, under select circumstances, the adjuvant composition may be designed to induce an immune response predominantly of the Th1 type or Th2 type. High levels of Th1-type cytokines (e.g., IFN-γ, TNFα, IL-2 and IL-12) tend to favor the induction of cell mediated immune responses to an administered antigen. In contrast, high levels of Th2-type cytokines (e.g., IL-4, IL-5, IL-6 and IL-10) tend to favor the induction of humoral immune responses. Following application of a vaccine as provided herein, a patient will support an immune response that includes Th1- and Th2-type responses. Within a preferred embodiment, in which a response is predominantly Th1-type, the level of Th1-type cytokines will increase to a greater extent than the level of Th2-type cytokines. The levels of these cytokines may be readily assessed using standard assays. For a review of the families of cytokines, see Mosmann and Coffman, Ann. Rev. Immunol. 7:145-173, 1989.

Preferred adjuvants for use in eliciting a predominantly Th1-type response include, for example, a combination of monophosphoryl lipid A, preferably 3de-O-acylated monophosphoryl lipid A (3D-MPL), together with an aluminum salt. MPL adjuvants are available from Corixa Corporation (Seattle, WA; see US Patent Nos. 4,436,727; 4,877,611; 4,866,034 and 4,912,094). CpG-containing oligonucleotides (in which the CpG dinucleotide is unmethylated) also induce a predominantly Th1 response. Such oligonucleotides are well known and are described, for example, in WO 96/02555 and WO 99/33488. Immunostimulatory DNA sequences are also described, for example, by Sato et al., Science 273:352, 1996. Another preferred adjuvant is a saponin, preferably QS21 (Aquila Biopharmaceuticals Inc., Framingham, MA), which may be used alone or in combination with other adjuvants. For example, an enhanced system involves the combination of a monophosphoryl lipid A and saponin derivative, such as the combination of QS21 and 3D-MPL as described in WO 94/00153, or a less reactogenic composition where the QS21 is quenched with cholesterol, as described in WO 96/33739. Other preferred formulations comprise an oil-in-water emulsion and tocopherol. A particularly potent adjuvant formulation involving OS21, 3D-MPL and tocopherol in an oil-in-water emulsion is described in WO 95/17210.

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Other preferred adjuvants include Montanide ISA 720 (Seppic, France), SAF (Chiron, California, United States), ISCOMS (CSL), MF-59 (Chiron), the SBAS series of adjuvants (e.g., SBAS-2 or SBAS-4, available from SmithKline Beecham, Rixensart, Belgium), Detox (Corixa Corporation; Seattle, WA), RC-529 (Corixa Corporation; Seattle, WA) and other aminoalkyl glucosaminide 4-phosphates (AGPs), such as those described in pending U.S. Patent Application Serial Nos. 08/853,826 and 09/074,720, the disclosures of which are incorporated herein by reference in their entireties.

Any vaccine provided herein may be prepared using well known methods that result in a combination of antigen, immunostimulant and a suitable carrier or excipient. The compositions described herein may be administered as part of a sustained release formulation (i.e., a formulation such as a capsule, sponge or gel (composed of polysaccharides, for example) that effects a slow release of compound following administration). Such formulations may generally be prepared using well known technology (see, e.g., Coombes et al., Vaccine 14:1429-1438, 1996) and

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administered by, for example, oral, rectal or subcutaneous implantation, or by implantation at the desired target site. Sustained-release formulations may contain a polypeptide, polynucleotide or antibody dispersed in a carrier matrix and/or contained within a reservoir surrounded by a rate controlling membrane.

Carriers for use within such formulations are biocompatible, and may also be biodegradable; preferably the formulation provides a relatively constant level of active component release. Such carriers include microparticles of poly(lactide-coglycolide), as well as polyacrylate, latex, starch, cellulose and dextran. Other delayed-release carriers include supramolecular biovectors, which comprise a non-liquid hydrophilic core (e.g., a cross-linked polysaccharide or oligosaccharide) and, optionally, an external layer comprising an amphiphilic compound, such as a phospholipid (see e.g., U.S. Patent No. 5,151,254 and PCT applications WO 94/20078, WO/94/23701 and WO 96/06638). The amount of active compound contained within a sustained release formulation depends upon the site of implantation, the rate and expected duration of release and the nature of the condition to be treated or prevented.

Any of a variety of delivery vehicles may be employed within pharmaceutical compositions and vaccines to facilitate production of an antigen-specific immune response that targets *Chlamydia*-infected cells. Delivery vehicles include antigen presenting cells (APCs), such as dendritic cells, macrophages, B cells, monocytes and other cells that may be engineered to be efficient APCs. Such cells may, but need not, be genetically modified to increase the capacity for presenting the antigen, to improve activation and/or maintenance of the T cell response, to have anti-*Chlamydia* effects *per se* and/or to be immunologically compatible with the receiver (*i.e.*, matched HLA haplotype). APCs may generally be isolated from any of a variety of biological fluids and organs, and may be autologous, allogeneic, syngeneic or xenogeneic cells.

Certain preferred embodiments of the present invention use dendritic cells or progenitors thereof as antigen-presenting cells. Dendritic cells are highly potent APCs (Banchereau and Steinman, *Nature 392*:245-251, 1998) and have been shown to be effective as a physiological adjuvant for eliciting prophylactic or therapeutic immunity (*see* Timmerman and Levy, *Ann. Rev. Med. 50*:507-529, 1999). In general, dendritic cells may be identified based on their typical shape (stellate *in situ*, with

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PCT/US01/23121

marked cytoplasmic processes (dendrites) visible *in vitro*), their ability to take up, process and present antigens with high efficiency, and their ability to activate naïve T cell responses. Dendritic cells may, of course, be engineered to express specific cell-surface receptors or ligands that are not commonly found on dendritic cells *in vivo* or *ex vivo*, and such modified dendritic cells are contemplated by the present invention. As an alternative to dendritic cells, secreted vesicles antigen-loaded dendritic cells (called exosomes) may be used within a vaccine (*see Zitvogel et al.*, *Nature Med. 4*:594-600, 1998).

Dendritic cells and progenitors may be obtained from peripheral blood, bone marrow, lymph nodes, spleen, skin, umbilical cord blood or any other suitable tissue or fluid. For example, dendritic cells may be differentiated *ex vivo* by adding a combination of cytokines such as GM-CSF, IL-4, IL-13 and/or TNFα to cultures of monocytes harvested from peripheral blood. Alternatively, CD34 positive cells harvested from peripheral blood, umbilical cord blood or bone marrow may be differentiated into dendritic cells by adding to the culture medium combinations of GM-CSF, IL-3, TNFα, CD40 ligand, LPS, flt3 ligand and/or other compound(s) that induce differentiation, maturation and proliferation of dendritic cells.

Dendritic cells are conveniently categorized as "immature" and "mature" cells, which allows a simple way to discriminate between two well characterized phenotypes. However, this nomenclature should not be construed to exclude all possible intermediate stages of differentiation. Immature dendritic cells are characterized as APC with a high capacity for antigen uptake and processing, which correlates with the high expression of Fcy receptor and mannose receptor. The mature phenotype is typically characterized by a lower expression of these markers, but a high expression of cell surface molecules responsible for T cell activation such as class I and class II MHC, adhesion molecules (e.g., CD54 and CD11) and costimulatory molecules (e.g., CD40, CD80, CD86 and 4-1BB).

APCs may generally be transfected with a polynucleotide encoding a Chlamydial protein (or portion or other variant thereof) such that the Chlamydial polypeptide, or an immunogenic portion thereof, is expressed on the cell surface. Such transfection may take place ex vivo, and a composition or vaccine comprising such transfected cells may then be used for therapeutic purposes, as described herein.

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PCT/US01/23121

Alternatively, a gene delivery vehicle that targets a dendritic or other antigen presenting cell may be administered to a patient, resulting in transfection that occurs in vivo. In vivo and ex vivo transfection of dendritic cells, for example, may generally be performed using any methods known in the art, such as those described in WO 97/24447, or the gene gun approach described by Mahvi et al., Immunology and cell Biology 75:456-460, 1997. Antigen loading of dendritic cells may be achieved by incubating dendritic cells or progenitor cells with the Chlamydial polypeptide, DNA (naked or within a plasmid vector) or RNA; or with antigen-expressing recombinant bacterium or viruses (e.g., vaccinia, fowlpox, adenovirus or lentivirus vectors). Prior to loading, the polypeptide may be covalently conjugated to an immunological partner that provides T cell help (e.g., a carrier molecule). Alternatively, a dendritic cell may be pulsed with a nonconjugated immunological partner, separately or in the presence of the polypeptide.

Routes and frequency of administration of pharmaceutical compositions and vaccines, as well as dosage, will vary from individual to individual. In general, the pharmaceutical compositions and vaccines may be administered by injection (e.g., intracutaneous, intramuscular, intravenous or subcutaneous), intranasally (e.g., by aspiration) or orally. Between 1 and 3 doses may be administered for a 1-36 week period. Preferably, 3 doses are administered, at intervals of 3-4 months, and booster vaccinations may be given periodically thereafter. Alternate protocols may be appropriate for individual patients. A suitable dose is an amount of polypeptide or DNA that, when administered as described above, is capable of raising an immune response in an immunized patient sufficient to protect the patient from Chlamydial infection for at least 1-2 years. In general, the amount of polypeptide present in a dose (or produced in situ by the DNA in a dose) ranges from about 1 pg to about 100 mg per kg of host, typically from about 10 pg to about 1 mg, and preferably from about 100 pg to about 1 µg. Suitable dose sizes will vary with the size of the patient, but will typically range from about 0.1 mL to about 5 mL.

While any suitable carrier known to those of ordinary skill in the art may be employed in the pharmaceutical compositions of this invention, the type of carrier will vary depending on the mode of administration. For parenteral administration, such as subcutaneous injection, the carrier preferably comprises water, saline, alcohol, a fat, a wax or a buffer. For oral administration, any of the above carriers or a solid carrier,

such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, sucrose, and magnesium carbonate, may be employed. Biodegradable microspheres (e.g., polylactic galactide) may also be employed as carriers for the pharmaceutical compositions of this invention. Suitable biodegradable microspheres are disclosed, for example, in U.S. Patent Nos. 4,897,268 and 5,075,109.

In general, an appropriate dosage and treatment regimen provides the active compound(s) in an amount sufficient to provide therapeutic and/or prophylactic benefit. Such a response can be monitored by establishing an improved clinical outcome in treated patients as compared to non-treated patients. Increases in preexisting immune responses to a *Chlamydial* protein generally correlate with an improved clinical outcome. Such immune responses may generally be evaluated using standard proliferation, cytotoxicity or cytokine assays, which may be performed using samples obtained from a patient before and after treatment.

In another aspect, the present invention provides methods for using the polypeptides described above to diagnose Chlamydial infection. In this aspect, methods are provided for detecting Chlamydial infection in a biological sample, using one or more of the above polypeptides, either alone or in combination. For clarity, the term "polypeptide" will be used when describing specific embodiments of the inventive diagnostic methods. However, it will be clear to one of skill in the art that the fusion proteins of the present invention may also be employed in such methods.

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As used herein, a "biological sample" is any antibody-containing sample obtained from a patient. Preferably, the sample is whole blood, sputum, serum, plasma, saliva, cerebrospinal fluid or urine. More preferably, the sample is a blood, serum or plasma sample obtained from a patient. The polypeptides are used in an assay, as described below, to determine the presence or absence of antibodies to the polypeptide(s) in the sample, relative to a predetermined cut-off value. The presence of such antibodies indicates previous sensitization to *Chlamydia* antigens which may be indicative of *Chlamydia*-infection.

In embodiments in which more than one polypeptide is employed, the polypeptides used are preferably complementary (i.e., one component polypeptide will tend to detect infection in samples where the infection would not be detected by another component polypeptide). Complementary polypeptides may generally be identified by

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using each polypeptide individually to evaluate serum samples obtained from a series of patients known to be infected with Chlamydia. After determining which samples test positive (as described below) with each polypeptide, combinations of two or more polypeptides may be formulated that are capable of detecting infection in most, or all, of the samples tested.

81

PCT/US01/23121

A variety of assay formats are known to those of ordinary skill in the art for using one or more polypeptides to detect antibodies in a sample. See, e.g., Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, 1988, which is incorporated herein by reference. In a preferred embodiment, the assay involves the use of polypeptide immobilized on a solid support to bind to and remove the antibody from the sample. The bound antibody may then be detected using a detection reagent that contains a reporter group. Suitable detection reagents include antibodies that bind to the antibody/polypeptide complex and free polypeptide labeled with a reporter group (e.g., in a semi-competitive assay). Alternatively, a competitive assay may be utilized, in which an antibody that binds to the polypeptide is labeled with a reporter group and allowed to bind to the immobilized antigen after incubation of the antigen with the sample. The extent to which components of the sample inhibit the binding of the labeled antibody to the polypeptide is indicative of the reactivity of the sample with the immobilized polypeptide.

The solid support may be any solid material known to those of ordinary skill in the art to which the antigen may be attached. For example, the solid support may be a test well in a microtiter plate, or a nitrocellulose or other suitable membrane. Alternatively, the support may be a bead or disc, such as glass, fiberglass, latex or a plastic material such as polystyrene or polyvinylchloride. The support may also be a magnetic particle or a fiber optic sensor, such as those disclosed, for example, in U.S. Patent No. 5,359,681.

The polypeptides may be bound to the solid support using a variety of techniques known to those of ordinary skill in the art. In the context of the present invention, the term "bound" refers to both noncovalent association, such as adsorption, and covalent attachment (which may be a direct linkage between the antigen and functional groups on the support or may be a linkage by way of a cross-linking agent). Binding by adsorption to a well in a microtiter plate or to a membrane is preferred. In

82

such cases, adsorption may be achieved by contacting the polypeptide, in a suitable buffer, with the solid support for a suitable amount of time. The contact time varies with temperature, but is typically between about 1 hour and 1 day. In general, contacting a well of a plastic microtiter plate (such as polystyrene or polyvinylchloride) with an amount of polypeptide ranging from about 10 ng to about 1 μ g, and preferably about 100 ng, is sufficient to bind an adequate amount of antigen.

Covalent attachment of polypeptide to a solid support may generally be achieved by first reacting the support with a bifunctional reagent that will react with both the support and a functional group, such as a hydroxyl or amino group, on the polypeptide. For example, the polypeptide may be bound to supports having an appropriate polymer coating using benzoquinone or by condensation of an aldehyde group on the support with an amine and an active hydrogen on the polypeptide (see, e.g., Pierce Immunotechnology Catalog and Handbook, 1991, at A12-A13).

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In certain embodiments, the assay is an enzyme linked immunosorbent assay (ELISA). This assay may be performed by first contacting a polypeptide antigen that has been immobilized on a solid support, commonly the well of a microtiter plate, with the sample, such that antibodies to the polypeptide within the sample are allowed to bind to the immobilized polypeptide. Unbound sample is then removed from the immobilized polypeptide and a detection reagent capable of binding to the immobilized antibody-polypeptide complex is added. The amount of detection reagent that remains bound to the solid support is then determined using a method appropriate for the specific detection reagent.

More specifically, once the polypeptide is immobilized on the support as described above, the remaining protein binding sites on the support are typically blocked. Any suitable blocking agent known to those of ordinary skill in the art, such as bovine serum albumin (BSA) or Tween 20TM (Sigma Chemical Co., St. Louis, MO) may be employed. The immobilized polypeptide is then incubated with the sample, and antibody is allowed to bind to the antigen. The sample may be diluted with a suitable dilutent, such as phosphate-buffered saline (PBS) prior to incubation. In general, an appropriate contact time (*i.e.*, incubation time) is that period of time that is sufficient to detect the presence of antibody within an HGE-infected sample. Preferably, the contact time is sufficient to achieve a level of binding that is at least 95% of that achieved at

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equilibrium between bound and unbound antibody. Those of ordinary skill in the art will recognize that the time necessary to achieve equilibrium may be readily determined by assaying the level of binding that occurs over a period of time. At room temperature, an incubation time of about 30 minutes is generally sufficient.

Unbound sample may then be removed by washing the solid support with an appropriate buffer, such as PBS containing 0.1% Tween 20™. Detection reagent may then be added to the solid support. An appropriate detection reagent is any compound that binds to the immobilized antibody-polypeptide complex and that can be detected by any of a variety of means known to those in the art. Preferably, the detection reagent contains a binding agent (such as, for example, Protein A, Protein G, immunoglobulin, lectin or free antigen) conjugated to a reporter group. Preferred reporter groups include enzymes (such as horseradish peroxidase), substrates, cofactors, inhibitors, dyes, radionuclides, luminescent groups, fluorescent groups and biotin. The conjugation of binding agent to reporter group may be achieved using standard methods known to those of ordinary skill in the art. Common binding agents may also be purchased conjugated to a variety of reporter groups from many commercial sources (e.g., Zymed Laboratories, San Francisco, CA, and Pierce, Rockford, IL).

The detection reagent is then incubated with the immobilized antibody-polypeptide complex for an amount of time sufficient to detect the bound antibody. An appropriate amount of time may generally be determined from the manufacturer's instructions or by assaying the level of binding that occurs over a period of time. Unbound detection reagent is then removed and bound detection reagent is detected using the reporter group. The method employed for detecting the reporter group depends upon the nature of the reporter group. For radioactive groups, scintillation counting or autoradiographic methods are generally appropriate. Spectroscopic methods may be used to detect dyes, luminescent groups and fluorescent groups. Biotin may be detected using avidin, coupled to a different reporter group (commonly a radioactive or fluorescent group or an enzyme). Enzyme reporter groups may generally be detected by the addition of substrate (generally for a specific period of time), followed by spectroscopic or other analysis of the reaction products.

To determine the presence or absence of anti-Chlamydia antibodies in the sample, the signal detected from the reporter group that remains bound to the solid

support is generally compared to a signal that corresponds to a predetermined cut-off value. In one preferred embodiment, the cut-off value is the average mean signal obtained when the immobilized antigen is incubated with samples from an uninfected patient. In general, a sample generating a signal that is three standard deviations above the predetermined cut-off value is considered positive for Chlamydia-infection. In an alternate preferred embodiment, the cut-off value is determined using a Receiver Operator Curve, according to the method of Sackett et al., Clinical Epidemiology: A Basic Science for Clinical Medicine, Little Brown and Co., 1985, pp. 106-107. Briefly, in this embodiment, the cut-off value may be determined from a plot of pairs of true positive rates (i.e., sensitivity) and false positive rates (100%-specificity) that correspond to each possible cut-off value for the diagnostic test result. The cut-off value on the plot that is the closest to the upper left-hand corner (i.e., the value that encloses the largest area) is the most accurate cut-off value, and a sample generating a signal that is higher than the cut-off value determined by this method may be considered positive. Alternatively, the cut-off value may be shifted to the left along the plot, to minimize the false positive rate, or to the right, to minimize the false negative rate. In general, a sample generating a signal that is higher than the cut-off value determined by this method is considered positive for Chlamydial infection.

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In a related embodiment, the assay is performed in a rapid flow-through or strip test format, wherein the antigen is immobilized on a membrane, such as nitrocellulose. In the flow-through test, antibodies within the sample bind to the immobilized polypeptide as the sample passes through the membrane. A detection reagent (e.g., protein A-colloidal gold) then binds to the antibody-polypeptide complex as the solution containing the detection reagent flows through the membrane. The detection of bound detection reagent may then be performed as described above. In the strip test format, one end of the membrane to which polypeptide is bound is immersed in a solution containing the sample. The sample migrates along the membrane through a region containing detection reagent and to the area of immobilized polypeptide. Concentration of detection reagent at the polypeptide indicates the presence of anti-Chlamydia antibodies in the sample. Typically, the concentration of detection reagent at that site generates a pattern, such as a line, that can be read visually. The absence of such a pattern indicates a negative result. In general, the amount of polypeptide

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immobilized on the membrane is selected to generate a visually discernible pattern when the biological sample contains a level of antibodies that would be sufficient to generate a positive signal in an ELISA, as discussed above. Preferably, the amount of polypeptide immobilized on the membrane ranges from about 25 ng to about 1 µg, and more preferably from about 50 ng to about 500 ng. Such tests can typically be performed with a very small amount (e.g., one drop) of patient serum or blood.

85

PCT/US01/23121

Of course, numerous other assay protocols exist that are suitable for use with the polypeptides of the present invention. The above descriptions are intended to be exemplary only. One example of an alternative assay protocol which may be usefully employed in such methods is a Western blot, wherein the proteins present in a biological sample are separated on a gel, prior to exposure to a binding agent. Such techniques are well known to those of skill in the art.

The present invention further provides agents, such as antibodies and antigen-binding fragments thereof, that specifically bind to a Chlamydial protein. As used herein, an antibody, or antigen-binding fragment thereof, is said to "specifically bind" to a Chlamydial protein if it reacts at a detectable level (within, for example, an ELISA) with a Chlamydial protein, and does not react detectably with unrelated proteins under similar conditions. As used herein, "binding" refers to a noncovalent association between two separate molecules such that a complex is formed. The ability to bind may be evaluated by, for example, determining a binding constant for the formation of the complex. The binding constant is the value obtained when the concentration of the complex is divided by the product of the component concentrations. In general, two compounds are said to "bind," in the context of the present invention, when the binding constant for complex formation exceeds about 10³ L/mol. The binding constant may be determined using methods well known in the art.

Binding agents may be further capable of differentiating between patients with and without a Chlamydial infection using the representative assays provided herein. In other words, antibodies or other binding agents that bind to a Chlamydial protein will generate a signal indicating the presence of a Chlamydial infection in at least about 20% of patients with the disease, and will generate a negative signal indicating the absence of the disease in at least about 90% of individuals without infection. To determine whether a binding agent satisfies this requirement, biological

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samples (e.g., blood, sera, sputum urine and/or tissue biopsies) from patients with and without Chlamydial infection (as determined using standard clinical tests) may be assayed as described herein for the presence of polypeptides that bind to the binding agent. It will be apparent that a statistically significant number of samples with and without the disease should be assayed. Each binding agent should satisfy the above criteria; however, those of ordinary skill in the art will recognize that binding agents may be used in combination to improve sensitivity.

86

PCT/US01/23121

Any agent that satisfies the above requirements may be a binding agent. For example, a binding agent may be a ribosome, with or without a peptide component, an RNA molecule or a polypeptide. In a preferred embodiment, a binding agent is an antibody or an antigen-binding fragment thereof. Antibodies may be prepared by any of a variety of techniques known to those of ordinary skill in the art. See, e.g., Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, 1988. In general, antibodies can be produced by cell culture techniques, including the generation of monoclonal antibodies as described herein, or via transfection of antibody genes into suitable bacterial or mammalian cell hosts, in order to allow for the production of recombinant antibodies. In one technique, an immunogen comprising the polypeptide is initially injected into any of a wide variety of mammals (e.g., mice, rats, rabbits, sheep or goats). In this step, the polypeptides of this invention may serve as the immunogen without modification. Alternatively, particularly for relatively short polypeptides, a superior immune response may be elicited if the polypeptide is joined to a carrier protein, such as bovine serum albumin or keyhole limpet hemocyanin. The immunogen is injected into the animal host, preferably according to a predetermined schedule incorporating one or more booster immunizations, and the animals are bled periodically. Polyclonal antibodies specific for the polypeptide may then be purified from such antisera by, for example, affinity chromatography using the polypeptide coupled to a suitable solid support.

Monoclonal antibodies specific for an antigenic polypeptide of interest may be prepared, for example, using the technique of Kohler and Milstein, Eur. J. Immunol. 6:511-519, 1976, and improvements thereto. Briefly, these methods involve the preparation of immortal cell lines capable of producing antibodies having the desired specificity (i.e., reactivity with the polypeptide of interest). Such cell lines may 87

WO 02/08267

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be produced, for example, from spleen cells obtained from an animal immunized as described above. The spleen cells are then immortalized by, for example, fusion with a myeloma cell fusion partner, preferably one that is syngeneic with the immunized animal. A variety of fusion techniques may be employed. For example, the spleen cells and myeloma cells may be combined with a nonionic detergent for a few minutes and then plated at low density on a selective medium that supports the growth of hybrid cells, but not myeloma cells. A preferred selection technique uses HAT (hypoxanthine, aminopterin, thymidine) selection. After a sufficient time, usually about 1 to 2 weeks, colonies of hybrids are observed. Single colonies are selected and their culture supernatants tested for binding activity against the polypeptide. Hybridomas having high reactivity and specificity are preferred.

PCT/US01/23121

Monoclonal antibodies may be isolated from the supernatants of growing hybridoma colonies. In addition, various techniques may be employed to enhance the yield, such as injection of the hybridoma cell line into the peritoneal cavity of a suitable vertebrate host, such as a mouse. Monoclonal antibodies may then be harvested from the ascites fluid or the blood. Contaminants may be removed from the antibodies by conventional techniques, such as chromatography, gel filtration, precipitation, and extraction. The polypeptides of this invention may be used in the purification process in, for example, an affinity chromatography step.

Within certain embodiments, the use of antigen-binding fragments of antibodies may be preferred. Such fragments include Fab fragments, which may be prepared using standard techniques. Briefly, immunoglobulins may be purified from rabbit serum by affinity chromatography on Protein A bead columns (Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, 1988) and digested by papain to yield Fab and Fc fragments. The Fab and Fc fragments may be separated by affinity chromatography on protein A bead columns.

Monoclonal antibodies of the present invention may be coupled to one or more therapeutic agents. Suitable agents in this regard include radionuclides, differentiation inducers, drugs, toxins, and derivatives thereof. Preferred radionuclides include ⁹⁰Y, ¹²³I, ¹²⁵I, ¹³¹I, ¹⁸⁶Re, ¹⁸⁸Re, ²¹¹At, and ²¹²Bi. Preferred drugs include methotrexate, and pyrimidine and purine analogs. Preferred differentiation inducers include phorbol esters and butyric acid. Preferred toxins include ricin, abrin, diptheria

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toxin, cholera toxin, gelonin, Pseudomonas exotoxin, Shigella toxin, and pokeweed antiviral protein.

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PCT/US01/23121

A therapeutic agent may be coupled (e.g., covalently bonded) to a suitable monoclonal antibody either directly or indirectly (e.g., via a linker group). A direct reaction between an agent and an antibody is possible when each possesses a substituent capable of reacting with the other. For example, a nucleophilic group, such as an amino or sulfhydryl group, on one may be capable of reacting with a carbonyl-containing group, such as an anhydride or an acid halide, or with an alkyl group containing a good leaving group (e.g., a halide) on the other.

Alternatively, it may be desirable to couple a therapeutic agent and an antibody via a linker group. A linker group can function as a spacer to distance an antibody from an agent in order to avoid interference with binding capabilities. A linker group can also serve to increase the chemical reactivity of a substituent on an agent or an antibody, and thus increase the coupling efficiency. An increase in chemical reactivity may also facilitate the use of agents, or functional groups on agents, which otherwise would not be possible.

It will be evident to those skilled in the art that a variety of bifunctional or polyfunctional reagents, both homo- and hetero-functional (such as those described in the catalog of the Pierce Chemical Co., Rockford, IL), may be employed as the linker group. Coupling may be effected, for example, through amino groups, carboxyl groups, sulfhydryl groups or oxidized carbohydrate residues. There are numerous references describing such methodology, e.g., U.S. Patent No. 4,671,958, to Rodwell et al.

Where a therapeutic agent is more potent when free from the antibody portion of the immunoconjugates of the present invention, it may be desirable to use a linker group which is cleavable during or upon internalization into a cell. A number of different cleavable linker groups have been described. The mechanisms for the intracellular release of an agent from these linker groups include cleavage by reduction of a disulfide bond (e.g., U.S. Patent No. 4,489,710, to Spitler), by irradiation of a photolabile bond (e.g., U.S. Patent No. 4,625,014, to Senter et al.), by hydrolysis of derivatized amino acid side chains (e.g., U.S. Patent No. 4,638,045, to Kohn et al.), by serum complement-mediated hydrolysis (e.g., U.S. Patent No. 4,569,789, to Blattler et al.).

It may be desirable to couple more than one agent to an antibody. In one embodiment, multiple molecules of an agent are coupled to one antibody molecule. In another embodiment, more than one type of agent may be coupled to one antibody. Regardless of the particular embodiment, immunoconjugates with more than one agent may be prepared in a variety of ways. For example, more than one agent may be coupled directly to an antibody molecule, or linkers which provide multiple sites for attachment can be used. Alternatively, a carrier can be used.

A carrier may bear the agents in a variety of ways, including covalent bonding either directly or via a linker group. Suitable carriers include proteins such as albumins (e.g., U.S. Patent No. 4,507,234, to Kato et al.), peptides and polysaccharides such as aminodextran (e.g., U.S. Patent No. 4,699,784, to Shih et al.). A carrier may also bear an agent by noncovalent bonding or by encapsulation, such as within a liposome vesicle (e.g., U.S. Patent Nos. 4,429,008 and 4,873,088). Carriers specific for radionuclide agents include radiohalogenated small molecules and chelating compounds. For example, U.S. Patent No. 4,735,792 discloses representative radiohalogenated small molecules and their synthesis. A radionuclide chelate may be formed from chelating compounds that include those containing nitrogen and sulfur atoms as the donor atoms for binding the metal, or metal oxide, radionuclide. For example, U.S. Patent No. 4,673,562, to Davison et al. discloses representative chelating compounds and their synthesis.

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A variety of routes of administration for the antibodies and immunoconjugates may be used. Typically, administration will be intravenous, intramuscular, subcutaneous or in site-specific regions by appropriate methods. It will be evident that the precise dose of the antibody/immunoconjugate will vary depending upon the antibody used, the antigen density, and the rate of clearance of the antibody.

Antibodies may be used in diagnostic tests to detect the presence of *Chlamydia* antigens using assays similar to those detailed above and other techniques well known to those of skill in the art, thereby providing a method for detecting Chlamydial infection in a patient.

Diagnostic reagents of the present invention may also comprise DNA sequences encoding one or more of the above polypeptides, or one or more portions thereof. For example, at least two oligonucleotide primers may be employed in a

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PCT/US01/23121

polymerase chain reaction (PCR) based assay to amplify *Chlamydia*-specific cDNA derived from a biological sample, wherein at least one of the oligonucleotide primers is specific for a DNA molecule encoding a polypeptide of the present invention. The presence of the amplified cDNA is then detected using techniques well known in the art, such as gel electrophoresis. Similarly, oligonucleotide probes specific for a DNA molecule encoding a polypeptide of the present invention may be used in a hybridization assay to detect the presence of an inventive polypeptide in a biological sample.

As used herein, the term "oligonucleotide primer/probe specific for a DNA molecule" means an oligonucleotide sequence that has at least about 80%, preferably at least about 90% and more preferably at least about 95%, identity to the DNA molecule in question. Oligonucleotide primers and/or probes which may be usefully employed in the inventive diagnostic methods preferably have at least about 10-40 nucleotides. In a preferred embodiment, the oligonucleotide primers comprise at least about 10 contiguous nucleotides of a DNA molecule encoding one of the polypeptides disclosed herein. Preferably, oligonucleotide probes for use in the inventive diagnostic methods comprise at least about 15 contiguous oligonucleotides of a DNA molecule encoding one of the polypeptides disclosed herein. Techniques for both PCR based assays and hybridization assays are well known in the art (see, for example, Mullis et al. Ibid; Ehrlich, Ibid). Primers or probes may thus be used to detect Chlamydia-specific sequences in biological samples. DNA probes or primers comprising oligonucleotide sequences described above may be used alone or in combination with each other.

The following Examples are offered by way of illustration and not by way of limitation.

EXAMPLE 1

ISOLATION OF DNA SEQUENCES ENCODING CHLAMYDIA ANTIGENS

Chlamydia antigens of the present invention were isolated by expression cloning of a genomic DNA library of Chlamydia trachomatis LGV II essentially as described by Sanderson et al. (J. Exp. Med., 1995, 182:1751-1757) and were shown to induce PBMC proliferation and IFN- γ in an immunoreactive T cell line.

91

A Chlamydia-specific T cell line was generated by stimulating PBMCs from a normal donor with no history of chlamydial genital tract infection with elementary bodies of Chlamydia trachomatis LGV II. This T cell line, referred to as TCL-8, was found to recognize both Chlamydia trachomatis and Chlamydia pneumonia infected monocyte-derived dendritic cells.

A randomly sheared genomic library of *Chlamydia trachomatis* LGV II was constructed in Lambda ZAP (Stratagene, La Jolla, CA) and the amplified library plated out in 96 well microtiter plates at a density of 30 clones/well. Bacteria were induced to express recombinant protein in the presence of 2 mM IPTG for 3 h, then pelleted and resuspended in 200 μl of RPMI 10% FBS. 10 μl of the induced bacterial suspension was transferred to 96 well plates containing autologous monocyte-derived dendritic cells. After a 2 h incubation, dendritic cells were washed to remove free *E. coli* and *Chlamydia*-specific T cells were added. Positive *E. coli* pools were identified by determining IFN-γ production and proliferation of the T cells in response to the pools.

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Four positive pools were identified, which were broken down to yield four pure clones (referred to as 1-B1-66, 4-D7-28, 3-G3-10 and 10-C10-31), with insert sizes of 481 bp, 183 bp, 110 bp and 1400 bp, respectively. The determined DNA sequences for 1-B1-66, 4-D7-28, 3-G3-10 and 10-C10-31 are provided in SEQ ID NO: 1-4, respectively. Clone 1-B1-66 is approximately in region 536690 of the C. trachomatis genome (NCBI C. trachomatis database). Within clone 1-B1-66, an open reading frame (ORF) has been identified (nucleotides 115 - 375) that encodes a previously identified 9 kDa protein (Stephens, et al. Genbank Accession No. AE001320), the sequence of which is provided in SEQ ID NO: 5). Clone 4-D7-28 is a smaller region of the same ORF (amino acids 22-82 of 1-B1-66). Clone 3-G3-10 is approximately in region 74559 of the C. trachomatis genome. The insert is cloned in the antisense orientation with respect to its orientation in the genome. The clone 10-C10-31 contains an open reading frame that corresponds to a previously published sequence for S13 ribosomal protein from Chlamydia trachomatis (Gu, L. et al. J. Bacteriology, 177:2594-2601, 1995). The predicted protein sequences for 4-D7-28 and 10-C10-31 are provided in SEQ ID NO: 6 and 12, respectively. Predicted protein sequences for 3-G3-10 are provided in SEQ ID NO: 7-11.

In a related series of screening studies, an additional T cell line was used to screen the genomic DNA library of *Chlamydia trachomatis* LGV II described above. A *Chlamydia*-specific T cell line (TCT-1) was derived from a patient with a chlamydial genital tract infection by stimulating patient PBMC with autologous monocyte-derived dendritic cells infected with elementary bodies of *Chlamydia trachomatis* LGV II. One clone, 4C9-18 (SEQ ID NO: 21), containing a 1256 bp insert, elicited a specific immune response, as measured by standard proliferation assays, from the *Chlamydia*-specific T cell line TCT-1. Subsequent analysis revealed this clone to contain three known sequences: lipoamide dehydrogenase (Genbank Accession No. AE001326), disclosed in SEQ ID NO: 22; a hypothetical protein CT429 (Genbank Accession No. AE001316), disclosed in SEQ ID NO: 23; and part of an open reading frame of ubiquinone methyltransferase CT428 (Genbank Accession No. AE001316), disclosed in SEQ ID NO: 24.

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In further studies involving clone 4C9-18 (SEQ ID NO: 21), the full-length amino acid sequence for lipoamide dehydrognase (SEQ ID NO: 22) from *C. trachomatis* (LGV II) was expressed in clone CtL2-LPDA-FL, as disclosed in SEQ ID NO: 90.

To further characterize the open reading frame containing the T cell stimulating epitope(s), a cDNA fragment containing nucleotides 1-695 of clone 4C9-18 with a cDNA sequence encoding a 6X-Histidine tag on the amino terminus was subcloned into the Ndel/EcoRI site of the pET17b vector (Novagen, Madison, WI), referred to as clone 4C9-18#2 BL21 pLysS (SEQ ID NO: 25, with the corresponding amino acid sequence provided in SEQ ID NO: 26) and transformed into *E. coli*. Selective induction of the transformed *E. coli* with 2 mM IPTG for three hours resulted in the expression of a 26 kDa protein from clone 4C9-18#2 BL21 pLysS, as evidenced by standard Coomassie-stained SDS-PAGE. To determine the immunogenicity of the protein encoded by clone 4C9-18#2 BL21 pLysS, *E. coli* expressing the 26 kDa protein were titered onto 1 x 10⁴ monocyte-derived dendritic cells and incubated for two hours. The dendritic cell cultures were washed and 2.5 x 10⁴ T cells (TCT-1) added and allowed to incubate for an additional 72 hours, at which time the level of IFN-γ in the culture supernatant was determined by ELISA. As shown in Fig. 1, the T-cell line TCT-1 was found to respond to induced cultures as measured by IFN-g, indicating a

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Chlamydia-specific T-cell response against the lipoamide dehydrogenase sequence. Similarly, the protein encoded by clone 4C9-18#2 BL21 pLysS was shown to stimulate the TCT-1 T-cell line by standard proliferation assays.

Subsequent studies to identify additional *Chlamydia trachomatis* antigens using the above-described CD4+ T-cell expression cloning technique yielded additional clones. The TCT-1 and TCL-8 *Chlamydia*-specific T-cell lines, as well as the TCP-21 T-cell line were utilized to screen the *Chlamydia trachomatis* LGVII genomic library. The TCP-21 T-cell line was derived from a patient having a humoral immune response to *Chlamydia pnuemoniae*. The TCT-1 cell line identified 37 positive pools, the TCT-3 cell line identified 41 positive pools and the TCP-21 cell line identified 2 positive pools. The following clones were derived from 10 of these positive pools. Clone 11-A3-93 (SEQ ID NO: 64), identified by the TCP-21 cell line, is a 1339 bp genomic fragment sharing homology to the HAD superfamily (CT103). The second insert in the same clone shares homology with the fab I gene (CT104) present on the complementary strand. Clone 11-C12-91 (SEQ ID NO: 63), identified using the TCP-21 cell line, has a 269 bp insert that is part of the OMP2 gene (CT443) and shares homology with the 60 kDa cysteine rich outer membrane protein of *C. pnuemoniae*.

Clone 11-G10-46, (SEQ ID NO: 62), identified using the TCT-3 cell line, contains a 688 bp insert that shares homology to the hypothetical protein CT610. Clone 11-G1-34, (SEQ ID NO: 61), identified using the TCT-3 cell line, has two partial open reading frames (ORF) with an insert size of 1215 bp. One ORF shares homology to the malate dehydrogenase gene (CT376), and the other ORF shares homology to the glycogen hydrolase gene (CT042). Clone 11-H3-68, (SEQ ID NO: 60), identified using the TCT-3 cell line, has two ORFs with a total insert size of 1180 bp. One partial ORF encodes the plasmid-encoded PGP6-D virulence protein while the second ORF is a complete ORF for the L1 ribosomal gene (CT318). Clone 11-H4-28, (SEQ ID NO: 59), identified using the TCT-3 cell line, has an insert size of 552 bp and is part of the ORF for the dnaK gene (CT396). Clone 12-B3-95, (SEQ ID NO: 58), identified using the TCT-1 cell line, has an insert size of 463 bp and is a part of the ORF for the lipoamide dehydrogenase gene (CT557). Clones 15-G1-89 and 12-B3-95 are identical, (SEQ ID NO: 55 and 58, respectively), identified using the TCT-1 cell line, has an insert size of 463 bp and is part of the ORF for the lipoamide dehydrogenase gene

PCT/US01/23121

(CT557). Clone 12-G3-83, (SEQ ID NO: 57), identified using the TCT-1 cell line, has an insert size of 1537 bp and has part of the ORF for the hypothetical protein CT622.

Clone 23-G7-68, (SEQ ID NO: 79), identified using the TCT-3 cell line. contains a 950 bp insert and contains a small part of the L11 ribosomal ORF, the entire ORF for L1 ribosomal protein and a part of the ORF for L10 ribosomal protein. In addition, this clone also identified the patient lines CT4, CT5, CT11, CT12, and CHH037. Clone 22-F8-91, (SEQ ID NO: 80), identified using the TCT-1 cell line, contains a 395 bp insert that contains a part of the pmpC ORF on the complementary strand of the clone. Clone 21-E8-95, (SEQ ID NO: 81), identified using the TCT-3 cell line, contains a 2,085 bp insert which contains part of CT613 ORF, the complete ORF for CT612, the complete ORF for CT611 and part of the ORF for CT610. Clone 19-F12-57, (SEQ ID NO: 82), identified using the TCT-3 cell line, contains a 405 bp insert which contains part of the CT 858 ORF and a small part of the recA ORF. Clone 19-F12-53, (SEQ ID NO: 83), identified using the TCT-3 cell line, contains a 379 bp insert that is part of the ORF for CT455 encoding glutamyl tRNA synthetase. Clone 19-A5-54, (SEQ ID NO: 84), identified using the TCT-3 cell line, contains a 715 bp insert that is part of the ORF3 (complementary strand of the clone) of the cryptic plasmid. Clone 17-E11-72, (SEQ ID NO: 85), identified using the TCT-1 cell line, contains a 476 bp insert that is part of the ORF for Opp 2 and pmpD. The pmpD region of this clone is covered by the pmpD region of clone 15-H2-76. Clone 17-C1-77, (SEO ID NO: 86), identified using the the patient cell lines CT3, CT1, CT4, and CT12, contains a 1551 bp insert that is part of the CT857 ORF, as well as part of the CT858 ORF. Clone 15-H2-76, (SEQ ID NO: 87), identified using the TCT-1 cell line, contains a 3,031 bp insert that contains a large part of the pmpD ORF, part of the CT089 ORF, as well as part of the ORF for SycE. Clone 15-A3-26, (SEQ ID NO: 88), contains a 976 bp insert that contains part of the ORF for CT858. Clone 17-G4-36, (SEQ ID NO: 267), identified using the patient lines CL8, TCT-10, CT1, CT5, CT13, and CHH037, contains a 680 bp insert that is in frame with beta-gal in the plasmid and shares homology to part of the ORF for DNA-directed RNA polymerase beta subunit (CT315 in SerD).

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Several of the clones described above share homology to various polymorphic membrane proteins. The genomic sequence of *Chlamydia trachomatis* contains a family of nine polymorphic membrane protein genes, referred to as pmp.

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These genes are designated pmpA, pmpB, pmpC, pmpD, pmpE, pmpF, pmpG, pmpH and pmpI. Proteins expressed from these genes are believed to be of biological relevance in generating a protective immune response to a *Chlamydial* infection. In particular, pmpC, pmpD, pmpE and pmpI contain predictable signal peptides, suggesting they are outer membrane proteins, and therefore, potential immunological targets.

Based on the Chlamydia trachomatis LGVII serovar sequence, primer pairs were designed to PCR amplify the full-length fragments of pmpC, pmpD, pmpE, pmpG, pmpH and pmpI. The resulting fragments were subcloned into the DNA vaccine vector JA4304 or JAL, which is JA4304 with a modified linker (SmithKline Beecham, London, England). Specifically, PmpC was subcloned into the JAL vector using the 5' oligo GAT AGG CGC GCC GCA ATC ATG AAA TTT ATG TCA GCT ACT GCT G and the 3' oligo CAG AAC GCG TTT AGA ATG TCA TAC GAG CAC CGC A, as provided in SEQ ID NO: 197 and 198, respectively. PCR amplification of the gene under conditions well known in the art and ligation into the 5' ASCI/3' MluI sites of the JAL vector was completed after inserting the short nucleotide sequence GCAATC (SEQ ID NO: 199) upstream of the ATG to create a Kozak-like sequence. The resulting expression vector contained the full-length pmpC gene comprising 5325 nucleotides (SEQ ID NO: 173) containing the hypothetical signal sequence, which encodes a 187 kD protein (SEQ ID NO: 179). The pmpD gene was subcloned into the JA4304 vaccine vector following PCR amplification of the gene using the following oligos: 5' oligo-TGC AAT CAT GAG TTC GCA GAA AGA TAT AAA AAG C (SEQ ID NO: 200) and 3' oligo- CAG AGC TAG CTT AAA AGA TCA ATC GCA ATC CAG TAT TC (SEQ ID NO: 201). The gene was ligated into the a 5' blunted HIII/3' MluI site of the JA4304 vaccine vector using standard techniques well known in the art. The CAATC (SEQ ID NO: 202) was inserted upstream of the ATG to create a Kozak-like sequence. This clone is unique in that the last threonine of the HindIII site is missing due to the blunting procedure, as is the last glycine of the Kozak-like sequence. The insert, a 4593 nucleotide fragment (SEQ ID NO: 172) is the full-length gene for pmpD containing the hypothetical signal sequence, which encodes a 161 kD protein (SEQ ID NO: 178). PmpE was subcloned into the JA4304 vector using the 5' oligo- TGC AAT CAT GAA AAA AGC GTT TTT CTT TTT C (SEQ ID NO: 203), and the 3' oligo- CAG AAC

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PCT/US01/23121

GCG TCT AGA ATC GCA GAG CAA TTT C (SEQ ID NO: 204). Following PCR amplification, the gene was ligated into the 5' blunted HIII/3' MluI site of JA4304. To facilitate this, a short nucleotide sequence, TGCAATC (SEQ ID NO: 293), was added upstream of the initiation codon for creating a Kozak-like sequence and reconstituting the HindIII site. The insert is the full-length pmpE gene (SEQ ID NO: 171) containing the hypothetical signal sequence. The pmpE gene encodes a 105 kD protein (SEQ ID NO: 177). The pmpG gene was PCR amplified using the 5' oligo- GTG CAA TCA TGA TTC CTC AAG GAA TTT ACG (SEQ ID NO: 205), and the 3' oligo- CAG AAC GCG TTT AGA ACC GGA CTT TAC TTC C (SEQ ID NO: 206) and subcloned into the JA4304 vector. Similar cloning strategies were followed for the pmpI and pmpK genes. In addition, primer pairs were designed to PCR amplify the full-length or overlapping fragments of the pmp genes, which were then subcloned for protein expression in the pET17b vector (Novagen, Madison, WI) and transfected into E. coli BL21 pLysS for expression and subsequent purification utilizing the histidine-nickel chromatographic methodology provided by Novagen. Several of the genes encoding the recombinant proteins, as described below, lack the native signal sequence to facilitate expression of the protein. Full-length protein expression of pmpC was accomplished through expression of two overlapping fragments, representing the amino and carboxy Subcloning of the pmpC-amino terminal portion, which lacks the signal termini. sequence, (SEQ ID NO: 187, with the corresponding amino acid sequence provided in SEQ ID NO: 195) used the 5' oligo- CAG ACA TAT GCA TCA CCA TCA CCA TCA CGA GGC GAG CTC GAT CCA AGA TC (SEQ ID NO: 207), and the 3' oligo- CAG AGG TAC CTC AGA TAG CAC TCT CTC CTA TTA AAG TAG G (SEQ ID NO: 208) into the 5' NdeI/3' KPN cloning site of the vector. The carboxy terminus portion of the gene, pmpC-carboxy terminal fragment (SEO ID NO: 186, with the corresponding amino acid sequence provided in SEQ ID NO: 194), was subcloned into the 5' Nhel/3' KPN cloning site of the expression vector using the following primers: 5' oligo- CAG AGC TAG CAT GCA TCA CCA TCA CGT TAA GAT TGA GAA CTT CTC TGG C (SEQ ID NO: 209), and 3' oligo- CAG AGG TAC CTT AGA ATG TCA TAC GAG CAC CGC AG (SEQ ID NO: 210). PmpD was also expressed as two overlapping proteins. The pmpD-amino terminal portion, which lacks the signal sequence, (SEQ ID NO: 185, with the corresponding amino acid sequence provided in

SEO ID NO: 193) contains the initiating codon of the pET17b and is expressed as a 80 kD protein. For protein expression and purification purposes, a six-histidine tag follows the initiation codon and is fused at the 28th amino acid (nucleotide 84) of the gene. The following primers were used, 5' oligo, CAG ACA TAT GCA TCA CCA TCA CCA TCA CGG GTT AGC (SEQ ID NO: 211), and the 3' oligo- CAG AGG TAC CTC AGC TCC TCC AGC ACA CTC TCT TC (SEQ ID NO: 212), to splice into the 5' NdeI/3' KPN cloning site of the vector. The pmpD-carboxy terminus portion (SEQ ID NO: 184) was expressed as a 92 kD protein (SEQ ID NO: 192). For expression and subsequent purification, an additional methionine, alanine and serine was included. which represent the initiation codon and the first two amino acids from the pET17b vector. A six-histidine tag downstream of the methionine, alanine and serine is fused at the 691st amino acid (nucleotide 2073) of the gene. The 5' oligo- CAG AGC TAG CCA TCA CCA TCA CCA TCA CGG TGC TAT TTC TTG CTT ACG TGG (SEO ID NO: 213) and the 3' oligo- CAG AGG TAC TTn AAA AGA TCA ATC GCA ATC CAG TAT TCG (SEQ ID NO: 214) were used to subclone the insert into the 5' NheJ/3' KPN cloning site of the expression vector. PmpE was expressed as a 106kD protein (SEQ ID NO: 183 with the corresponding amino acid sequence provided in SEQ ID NO: 191). The pmpE insert also lacks the native signal sequence. PCR amplification of the gene under conditions well known in the art was performed using the following oligo primers: 5' oligo- CAG AGG ATC CAC ATC ACC ATC ACC ATC ACG GAC TAG CTA GAG AGG TTC (SEQ ID NO: 215), and the 3' oligo- CAG AGA ATT CCT AGA ATC GCA GAG CAA TTT C (SEQ ID NO: 216), and the amplified insert was ligated into a 5' BamHI/3' EcoRI site of JA4304. The short nucleotide sequence, as provided in SEQ ID NO: 217, was inserted upstream of the initiation codon for creating the Kozak-like sequence and reconstituting the HindIII site. The expressed protein contains the initiation codon and the downstream 21 amino acids from the pET17b expression vector, i.e., MASMTGGQQMGRDSSLVPSSDP (SEQ ID NO: 218). In addition, a six-histidine tag is included upstream of the sequence described above and is fused at the 28th amino acid (nucleotide 84) of the gene, which eliminates the hypothetical signal peptide. The sequences provided in SEQ ID NO: 183 with the corresponding amino acid sequence provided in SEQ ID NO: 191 do not include these additional sequences. The pmpG gene (SEQ ID NO: 182, with the corresponding

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amino acid sequence provided in SEQ ID No; 190) was PCR amplified under conditions well known in the art using the following oligo primers: 5' oligo- CAG AGG TAC CGC ATC ACC ATC ACC ATC ACA TGA TTC CTC AAG GAA TTT ACG (SEO ID NO: 219), and the 3' oligo- CAG AGC GGC CGC TTA GAA CCG GAC TTT ACT TCC (SEQ ID NO: 220), and ligated into the 5' KPN/3' NotI cloning site of the expression vector. The expressed protein contains an additional amino acid sequence at the amino end, namely, MASMTGGQQNGRDSSLVPHHHHHHH (SEQ ID NO: 221), which comprises the initiation codon and additional sequence from the pET17b expression vector. The pmpI gene (SEQ ID NO: 181, with the corresponding amino acid sequence provided in SEQ ID No; 189) was PCR amplified under conditions well known in the art using the following oligo primers: 5' oligo- CAG AGC TAG CCA TCA CCA TCA CCT CTT TGG CCA GGA TCC C (SEQ ID NO: 222), and the 3' oligo- CAG AAC TAG TCT AGA ACC TGT AAG TGG TCC (SEQ ID NO: 223), and ligted into the expression vector at the 5' Nhel/3' Spel cloning site. The 95 kD expressed protein contains the initiation codon plus an additional alanine and serine from the pET17b vector at the amino end of the protein. In addition, a six-histidine tag is fused at the 21st amino acid of the gene, which eliminates the hypothetical signal peptide.

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Clone 14H1-4, (SEQ ID NO: 56), identified using the TCT-3 cell line, contains a complete ORF for the TSA gene, thiol specific antioxidant — CT603 (the CT603 ORF is a homolog of CPn0778 from *C. pnuemoniae*). The TSA open reading frame in clone 14-H1-4 was amplified such that the expressed protein possess an additional methionine and a 6x histidine tag (amino terminal end). This amplified insert was sub-cloned into the Nde/EcoRI sites of the pET17b vector. Upon induction of this clone with IPTG, a 22.6 kDa protein was purified by Ni-NTA agarose affinity chromatography. The determined amino acid sequence for the 195 amino acid ORF of clone 14-H1-4 encoding the TSA gene is provided in SEQ ID NO: 65. Further analysis yielded a full-length clone for the TSA gene, referred to as CTL2-TSA-FL, with the full-length amino acid sequence provided in SEQ ID NO: 92.

Further studies yielded 10 additional clones identified by the TCT-1 and TCT-3 T-cell lines, as described above. The clones identified by the TCT-1 line are: 16-D4-22, 17-C5-19, 18-C5-2, 20-G3-45 and 21-C7-66; clones identified by the TCT-3

cell line are: 17-C10-31, 17-E2-9, 22-A1-49 and 22-B3-53. Clone 21-G12-60 was recognized by both the TCT-1 and TCT-3 T cell lines. In addition, clone 20-G3-45, which contained sequence specific for pmpB, was identified against the patient lines CT1 and CT4. Clone 16-D4-22 (SEQ ID NO: 119), identified using the TCT-1 cell line contains a 953 bp insert that contains two genes, parts of open reading frame 3 (ORF3) and ORF4 of the C. trachomatis plasmid for growth within mammalian cells. Clone 17-C5-19 (SEQ ID NO: 118), contains a 951 bp insert that contains part of the ORF for DT431, encoding for clpP 1 protease and part of the ORF for CT430 (diaminopimelate epimerase). Clone 18-C5-2 (SEQ ID NO: 117) is part of the ORF for S1 ribosomal protein with a 446 bp insert that was identified using the TCT-1 cell line. Clone 20-G3-45 (SEQ ID NO: 116), identified by the TCT-1 cell line, contains a 437 bp insert that is part of the pmpB gene (CT413). Clone 21-C7-8 (SEQ ID NO: 115), identified by the TCT-1 line, contains a 995bp insert that encodes part of the dnaK like protein. The insert of this clone does not overlap with the insert of the TCT-3 clone 11-H4-28 (SEO ID NO: 59), which was shown to be part of the dnaK gene CT396. Clone 17-C10-31 (SEQ ID NO: 114), identified by the TCT-3 cell line, contains a 976 bp insert. This clone contains part of the ORF for CT858, a protease containing IRBP and DHR domains. Clone 17-E2-9 (SEQ ID NO: 113) contains part of ORFs for two genes. CT611 and CT610, that span a 1142 bp insert. Clone 22-A1-49 (SEO ID NO: 112), identified using the TCT-3 line, also contains two genes in a 698 bp insert. Part of the ORF for CT660 (DNA gyrase{gyrA 2}) is present on the top strand where as the complete ORF for a hypothetical protein CT659 is present on the complementary strand. Clone 22-B3-53 (SEQ ID NO: 111), identified by the TCT-1 line, has a 267 bp insert that encodes part of the ORF for GroEL (CT110). Clone 21-G12-60 (SEQ ID NO: 110), identified by both the TCT-1 and TCT-3 cell lines contains a 1461 bp insert that contains partial ORFs for hypothetical proteins CT875, CT229 and CT228.

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Additional *Chlamydia* antigens were obtained by screening a genomic expression library of *Chlamydia trachomatis* (LGV II serovar) in Lambda Screen-1 vector (Novagen, Madison, WI) with sera pooled from several *Chlamydia*-infected individuals using techniques well known in the art. The following immuno-reactive clones were identified and the inserts containing *Chlamydia* genes sequenced: CTL2#1 (SEQ ID NO: 71); CTL2#2 (SEQ ID NO: 70); CTL2#3-5' (SEQ ID NO: 72, a first

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determined genomic sequence representing the 5' end); CTL2#3-3' (SEQ ID NO: 73, a second determined genomic sequence representing the 3' end); CTL2#4 (SEQ ID NO: 53); CTL2#5 (SEQ ID NO: 69); CTL2#6 (SEQ ID NO: 68); CTL2#7 (SEQ ID NO: 67); CTL2#8b (SEQ ID NO: 54); CTL2#9 (SEQ ID NO: 66); CTL2#10-5' (SEQ ID NO: 74, a first determined genomic sequence representing the 5' end); CTL2#10-3' (SEQ ID NO: 75, a second determined genomic sequence representing the 3' end); CTL2#11-5' (SEQ ID NO: 45, a first determined genomic sequence representing the 5' end); CTL2#11-3' (SEQ ID NO: 44, a second determined genomic sequence representing the 3' end); CTL2#12 (SEQ ID NO: 46); CTL2#16-5' (SEQ ID NO: 47); CTL2#18-5' (SEQ ID NO: 49, a first determined genomic sequence representing the 5' end); CTL2#18-3' (SEQ ID NO: 48, a second determined genomic sequence representing the 3' end); CTL2#19-5' (SEQ ID NO: 76, the determined genomic sequence representing the 5' end); CTL2#21 (SEQ ID NO: 50); CTL2#23 (SEQ ID NO: 51; and CTL2#24 (SEQ ID NO: 52).

Additional Chlamydia trachomatis antigens were identified serological expression cloning. These studies used sera pooled from several Chlamydia-infected individuals, as described above, but, IgA, and IgM antibodies were used in addition to IgG as a secondary antibody. Clones screened by this method enhance detection of antigens recognized by an early immune response to a Chlamydial infection, that is a mucosal humoral immune response. The following immunoreactive clones were characterized and the inserts containing Chlamydia genes sequenced: CTL2gam-1 (SEQ ID NO: 290), CTL2gam-2 (SEQ ID NO: 289), CTL2gam-5 (SEQ ID NO: 288), CTL2gam-6-3' (SEQ ID NO: 287, a second determined genomic sequence representing the 3' end), CTL2gam-6-5' (SEQ ID NO: 286, a first determined genomic sequence representing the 5' end), CTL2gam-8 (SEQ ID NO: 285), CTL2gam-10 (SEQ ID NO: 284), CTL2gam-13 (SEQ ID NO: 283), CTL2gam-15-3' (SEQ ID NO: 282, a second determined genomic sequence representing the 3' end), CTL2gam-15-5' (SEQ ID NO: 281, a first determined genomic sequence representing the 5' end), CTL2gam-17 (SEQ ID NO: 280), CTL2gam-18 (SEQ ID NO: 279), CTL2gam-21 (SEQ ID NO: 278), CTL2gam-23 (SEQ ID NO: 277), CTL2gam-24 (SEQ ID NO: 276), CTL2gam-26 (SEQ ID NO: 275), CTL2gam-27 (SEQ ID NO: 274), CTL2gam-28 (SEQ ID NO: 273), CTL2gam-30-3' (SEQ ID NO: 272, a second determined genomic sequence

101

representing the 3' end) and CTL2gam-30-5' (SEQ ID NO: 271, a first determined genomic sequence representing the 5' end).

EXAMPLE 2

INDUCTION OF T CELL PROLIFERATION AND INTERFERON-γ PRODUCTION BY CHLAMYDIA TRACHOMATIS ANTIGENS

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The ability of recombinant *Chlamydia trachomatis* antigens to induce T cell proliferation and interferon-γ production is determined as follows.

Proteins are induced by IPTG and purified by Ni-NTA agarose affinity chromatograph (Webb et al., *J. Immunology 157*:5034-5041, 1996). The purified polypeptides are then screened for the ability to induce T-cell proliferation in PBMC preparations. PBMCs from *C. trachomatis* patients as well as from normal donors whose T-cells are known to proliferate in response to *Chlamydia* antigens, are cultured in medium comprising RPMI 1640 supplemented with 10% pooled human serum and 50 μg/ml gentamicin. Purified polypeptides are added in duplicate at concentrations of 0.5 to 10 μg/mL. After six days of culture in 96-well round-bottom plates in a volume of 200 μl, 50 μl of medium is removed from each well for determination of IFN-γ levels, as described below. The plates are then pulsed with 1 μCi/well of tritiated thymidine for a further 18 hours, harvested and tritium uptake determined using a gas scintillation counter. Fractions that result in proliferation in both replicates three fold greater than the proliferation observed in cells cultured in medium alone are considered positive.

IFN-γ is measured using an enzyme-linked immunosorbent assay (ELISA). ELISA plates are coated with a mouse monoclonal antibody directed to human IFN-γ (PharMingen, San Diego, CA) in PBS for four hours at room temperature. Wells are then blocked with PBS containing 5% (W/V) non-fat dried milk for 1 hour at room temperature. The plates are washed six times in PBS/0.2% TWEEN-20 and samples diluted 1:2 in culture medium in the ELISA plates are incubated overnight at room temperature. The plates are again washed and a polyclonal rabbit anti-human IFN-γ serum diluted 1:3000 in PBS/10% normal goat serum is added to each well. The

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PCT/US01/23121

102

plates are then incubated for two hours at room temperature, washed and horseradish peroxidase-coupled anti-rabbit IgG (Sigma Chemical So., St. Louis, MO) is added at a 1:2000 dilution in PBS/5% non-fat dried milk. After a further two hour incubation at room temperature, the plates are washed and TMB substrate added. The reaction is stopped after 20 min with 1 N sulfuric acid. Optical density is determined at 450 nm using 570 nm as a reference wavelength. Fractions that result in both replicates giving an OD two fold greater than the mean OD from cells cultured in medium alone, plus 3 standard deviations, are considered positive.

Using the above methodology, recombinant 1B1-66 protein (SEQ ID NO: 5) as well as two synthetic peptides corresponding to amino acid residues 48-67 (SEQ ID NO: 13; referred to as 1-B1-66/48-67) and 58-77 (SEQ ID NO: 14, referred to as 1B1-66/58-77), respectively, of SEQ ID NO: 5, were found to induce a proliferative response and IFN-γ production in a Chlamydia-specific T cell line used to screen a genomic library of *C. trachomatis* LGV II.

Further studies have identified a *C. trachomatis*-specific T-cell epitope in the ribosomal S13 protein. Employing standard epitope mapping techniques well known in the art, two T-cell epitopes in the ribosomal S13 protein (rS13) were identified with a *Chlamydia*-specific T-cell line from donor CL-8 (T-cell line TCL-8 EB/DC). Fig. 8 illustrates that the first peptide, rS13 1-20 (SEQ ID NO: 106), is 100% identical with the corresponding *C. pneumoniae* sequence, explaining the cross-reactivity of the T-cell line to recombinant *C. trachomatis*- and *C. pneumoniae*-rS13. The response to the second peptide rS13 56-75 (SEQ ID NO: 108) is *C. trachomatis*-specific, indicating that the rS13 response in this healthy asymptomatic donor was elicited by exposure to *C. trachomatis* and not to *C. pneumoniae*, or any other microbial infection.

As described in Example 1, Clone 11-C12-91 (SEQ ID NO: 63), identified using the TCP-21 cell line, has a 269 bp insert that is part of the OMP2 gene (CT443) and shares homology with the 60 kDa cysteine rich outer membrane protein of *C. pneumoniae*, referred to as OMCB. To further define the reactive epitope(s), epitope mapping was performed using a series of overlapping peptides and the immunoassay previously described. Briefly, proliferative responses were determined by stimulating 2.5 x 10⁴ TCP-21 T-cells in the presence of 1 x 10⁴ monocyte-derived dendritic cells

103 -

with either non-infectious elementary bodies derived from *C. trachomatis* and *C. pneumoniae*, or peptides derived from the protein sequence of *C. trachomatis* or *C. pneumoniae* OMCB protein (0.1 µg/ml). The TCP-21 T-cells responded to epitopes CT-OMCB #167-186, CT-OMCB #171-190, CT-OMCB #171-186, and to a lesser extent, CT-OMCB #175-186 (SEQ ID NO: 249-252, respectively). Notably, the TCP-21 T-cell line also gave a proliferative response to the homologous *C. pneumoniae* peptide CP-OMCB #171-186 (SEQ ID NO: 253), which was equal to or greater than the response to the *C. trachomatis* peptides. The amino acid substitutions in position two (i.e., Asp for Glu) and position four (i.e., Cys for Ser) did not alter the proliferative response of the T-cells and therefore demonstrating this epitope to be a cross-reactive epitope between *C. trachomatis* and *C. pneumoniae*.

To further define the epitope described above, an additional T-cell line, TCT-3, was used in epitope mapping experiments. The immunoassays were performed as described above, except that only peptides from *C. trachomatis* were tested. The T-cells gave a proliferative response to two peptides, CT-OMCB #152-171 and CT-OMCB #157-176 (SEQ ID NO: 246 and 247, respectively), thereby defining an additional immunogenic epitope in the cysteine rich outer membrane protein of *C. trachomatis*.

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Clone 14H1-4, (SEQ ID NO: 56, with the corresponding full-length amino acid sequence provided in SEQ ID NO: 92), was identified using the TCT-3 cell line in the CD4 T-cell expression cloning system previously described, and was shown to contain a complete ORF for the, thiol specific antioxidant gene (CT603), referred to as TSA. Epitope mapping immunoassays were performed, as described above, to further define the epitope. The TCT-3 T-cells line exhibited a strong proliferative response to the overlapping peptides CT-TSA #96-115, CT-TSA #101-120 and CT-TSA #106-125 (SEQ ID NO: 254-256, respectively) demonstrating an immunoreactive epitope in the thiol specific antioxidant gene of *C. trachomatis* serovar LGVII.

104

EXAMPLE 3 PREPARATION OF SYNTHETIC POLYPEPTIDES

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Polypeptides may be synthesized on a Millipore 9050 peptide synthesizer using FMOC chemistry with HPTU (O-Benzotriazole-N,N,N',N'-tetramethyluronium hexafluorophosphate) activation. A Gly-Cys-Gly sequence may be attached to the amino terminus of the peptide to provide a method of conjugating or labeling of the peptide. Cleavage of the peptides from the solid support may be carried out using the following cleavage mixture: trifluoroacetic acid:ethanedithiol:thioanisole:water:phenol (40:1:2:2:3). After cleaving for 2 hours, the peptides may be precipitated in cold methyl-t-butyl-ether. The peptide pellets may then be dissolved in water containing 0.1% trifluoroacetic acid (TFA) and lyophilized prior to purification by C18 reverse phase HPLC. A gradient of 0-60% acetonitrile (containing 0.1% TFA) in water (containing 0.1% TFA) may be used to elute the peptides. Following lyophilization of the pure fractions, the peptides may be characterized using electrospray mass spectrometry and by amino acid analysis.

EXAMPLE 4

20 <u>ISOLATION AND CHARACTERIZATION OF DNA SEQUENCES ENCODING</u> <u>CHLAMYDIA ANTIGENS USING RETROVIRAL EXPRESSION VECTOR</u> SYSTEMS AND SUBSEQUENT IMMUNOLOGICAL ANALYSIS

A genomic library of *Chlamydia trachomatis* LGV II was constructed by limited digests using BamHI, BglII, BstYi and MboI restriction enzymes. The restriction digest fragments were subsequently ligated into the BamHI site of the retroviral vectors pBIB-KS1,2,3. This vector set was modified to contain a Kosak translation initiation site and stop codons in order to allow expression of proteins from short DNA genomic fragments, as shown in Fig. 2. DNA pools of 80 clones were prepared and transfected into the retroviral packaging line Phoenix-Ampho, as described in Pear, W.S., Scott, M.L. and Nolan, G.P., Generation of High Titre, Helperfree Retroviruses by Transient Transfection. Methods in Molecular Medicine: Gene

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Therapy Protocols, Humana Press, Totowa, NJ, pp. 41-57. The *Chlamydia* library in retroviral form was then transduced into H2-Ld expressing P815 cells, which were then used as target cells to stimulate an antigen specific T-cell line.

105

PCT/US01/23121

A Chlamydia-specific, murine H2^d restricted CD8+ T-cell line was expanded in culture by repeated rounds of stimulation with irradiated C. trachomatisinfected J774 cells and irradiated syngeneic spleen cells, as described by Starnbach, M., in J. Immunol., 153:5183, 1994. This Chlamydia-specific T-cell line was used to screen the above Chlamydia genomic library expressed by the retrovirally-transduced P815 cells. Positive DNA pools were identified by detection of IFN-y production using Elispot analysis (SEE Lalvani et al., J. Experimental Medicine 186:859-865, 1997).

Two positive pools, referred to as 2C7 and 2E10, were identified by IFNy Elispot assays. Stable transductants of P815 cells from pool 2C7 were cloned by limiting dilution and individual clones were selected based upon their capacity to elicit IFN-y production from the *Chlamydia*-specific CTL line. From this screening process, four positive clones were selected, referred to as 2C7-8, 2C7-9, 2C7-19 and 2C7-21. Similarly, the positive pool 2E10 was further screened, resulting in an additional positive clone, which contains three inserts. The three inserts are fragments of the CT016, tRNA syntase and clpX genes (SEQ ID NO: 268-270, respectively).

Transgenic DNA from these four positive 2C7 clones were PCR amplified using pBIB-KS specific primers to selectively amplify the Chlamydia DNA insert. Amplified inserts were gel purified and sequenced. One immunoreactive clone, 2C7-8 (SEQ ID NO: 15, with the predicted amino acid sequence provided in SEQ ID NO: 32), is a 160 bp fragment with homology to nucleotides 597304-597145 of Chlamydia trachomatis, serovar D (NCBI, BLASTN search; SEQ ID NO: 33, with the predicted amino acid sequence provided in SEQ ID NO: 34). The sequence of clone 2C7-8 maps within two putative open reading frames from the region of high homology described immediately above, and in particular, one of these putative open reading frames, consisting of a 298 amino acid fragment (SEQ ID NO: 16, with the predicted amino acid sequence provided in SEQ ID NO: 17), was demonstrated to exhibit immunological activity.

Full-length cloning of the 298 amino acid fragment (referred to as CT529 and/or the Cap1 gene) from serovar L2 was obtained by PCR amplification using 5'-

ttttgaagcaggtaggtgaatatg (forward) (SEQ ID NO: 159) and 5'-ttaagaaatttaaaaaatccctta (reverse) (SEQ ID NO: 160) primers, using purified *C. trachomatis* L2 genomic DNA as template. This PCR product was gel-purified, cloned into pCRBlunt (Invitrogen, Carlsbad, CA) for sequencing, and then subcloned into the *Eco*RI site of pBIB-KMS, a derivative of pBIB-KS for expression. The *Chlamydia pnuemoniae* homlogue of CT529 is provided in SEQ ID NO: 291, with the corresponding amino acid sequence provided in SEQ ID NO: 292.

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Full-length DNA encoding various CT529 serovars were amplified by PCR from bacterial lysates containing 10⁵ IFU, essentially as described (Denamur, E., C. Sayada, A. Souriau, J. Orfila, A. Rodolakis and J. Elion. 1991. J. Gen. Microbiol. 137: 2525). The following serovars were amplified as described: Ba (SEO ID NO: 134, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 135); E (BOUR) and E (MTW447) (SEQ ID NO: 122, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 123); F (NI1) (SEQ ID NO: 128, with the corresponding predicted amino acid sequence provided in SEO ID NO: 129); G; (SEO ID NO: 126, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 127); Ia (SEQ ID NO: 124, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 125); L1 (SEQ ID NO: 130, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 131); L3 (SEQ ID NO: 132, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 133); I (SEQ ID NO: 263, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 264); K (SEQ ID NO: 265, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 266); and MoPn (SEQ ID NO: 136, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 137). PCR reactions were performed with Advantage Genomic PCR Kit (Clontech, Palo Alto, CA) using primers specific for serovar L2 DNA (external to the ORF). Primers sequences were 5'ggtataatatctctctaaattttg (forward-SEQ ID NO: 161) and 5'-agataaaaaaggctgtttc' (reverse-SEQ ID NO: 162) except for MoPn which required 5'-ttttgaagcaggtaggtgaatatg (forward-SEQ ID NO: 163) and 5'-tttacaataagaaaagctaagcactttgt (reverse-SEQ ID NO: 164). PCR amplified DNA was purified with QIAquick PCR purification kit (Oiagen, Valencia, CA) and cloned in pCR2.1 (Invitrogen, Carlsbad, CA) for sequencing.

Sequencing of DNA derived from PCR amplified inserts of immunoreactive clones was done on an automated sequencer (ABI 377) using both a pBIB-KS specific forward primer 5'-ccttacacagtcctgctgac (SEQ ID NO: 165) and a reverse primer 3'-gtttccgggccctcacattg (SEQ ID NO: 166). PCRBlunt cloned DNA coding for CT529 serovar L2 and pCR2.1 cloned DNA coding for CT529 serovar Ba, E (BOUR), E (MTW447), F (NI1), G, Ia, K, L1, L3 and MoPn were sequenced using T7 promoter primer and universal M13 forward and M13 reverse primers.

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To determine if these two putative open reading frames (SEQ ID NO: 16 and 20) encoded a protein with an associated immunological function, overlapping peptides (17-20 amino acid lengths) spanning the lengths of the two open reading frames were synthesized, as described in Example 3. A standard chromium release assay was utilized to determine the percent specific lysis of peptide-pulsed H2^d restricted target cells. In this assay, aliquots of P815 cells (H2^d) were labeled at 37° C for one hour with 100 μ Ci of ⁵¹Cr in the presence or absence of 1 μ g/ml of the indicated peptides. Following this incubation, labeled P815 cells were washed to remove excess ⁵¹Cr and peptide, and subsequently plated in duplicate in microculture plates at a concentration of 1,000 cells/well. Effector CTL (Chlamydia-specific CD8 T cells) were added at the indicated effector:target ratios. Following a 4 hour incubation, supernatants were harvested and measured by gamma-counter for release of ⁵¹Cr into the supernatant. Two overlapping peptides from the 298 amino acid open reading frame did specifically stimulate the CTL line. The peptides represented in SEQ ID NO: 138-156 were synthesized, representing the translation of the L2 homologue of the serovar D open reading frame for CT529 (Cap1 gene) and 216 amino acid open reading frame. As shown in Fig. 3, peptides CtC7.8-12 (SEQ ID NO: 18, also referred to as Cap1#132-147, SEQ ID NO: 139) and CtC7.8-13 (SEQ ID NO: 19, also referred to as Cap1#138-155, SEQ ID NO: 140) were able to elicit 38 to 52% specific lysis, respectively, at an effector to target ratio of 10:1. Notably, the overlap between these two peptides contained a predicted H2^d (K^d and L^d) binding peptide. A 10 amino acid peptide was synthesized to correspond to this overlapping sequence (SEQ ID NO: 31) and was found to generate a strong immune response from the anti-Chlamydia CTL line by elispot assay. Significantly, a search of the most recent Genbank database revealed no proteins have previously been described for this gene. Therefore, the putative open 5

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WO 02/08267 PCT/US01/23121

reading frame encoding clone 2C7-8 (SEQ ID NO: 15) defines a gene which encompasses an antigen from *Chlamydia* capable of stimulating antigen-specific CD8+ T-cells in a MHC-I restricted manner, demonstrating this antigen could be used to develop a vaccine against *Chlamydia*.

108

To confirm these results and to further map the epitope, truncated peptides (SEQ ID NO: 138-156) were made and tested for recognition by the T-cells in an IFN-g ELISPOT assay. Truncations of either Ser139 (Cap1#140-147, SEQ ID NO: 146) or Leu147 (Cap1#138-146, SEQ ID NO: 147) abrogate T-cell recognition. These results indicate that the 9-mer peptide Cap1#139-147 (SFIGGITYL, SEQ ID NO: 145) is the minimal epitope recognized by the *Chlamydia*-specific T-cells.

Sequence alignments of Cap1 (CT529) from selected serovars of *C. trachomatis* (SEQ ID NO: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139) shows one of the amino acid differences is found in position 2 of the proposed epitope. The homologous serovar D peptide is SIIGGITYL (SEQ ID NO: 168). The ability of SFIGGITYL and SIIGGITYL to target cells for recognition by the *Chlamydia* specific T-cells was compared. Serial dilutions of each peptide were incubated with P815 cells and tested for recognition by the T-cells in a ⁵¹Cr release assay, as described above. The *Chlamydia*-specific T-cells recognize the serovar L2 peptide at a minimum concentration of 1 nM and the serovar D peptide at a minimum concentration of 10 nM.

Further studies have shown that a Cap1#139-147-specific T-cell clone recognizes *C. trachomatis* infected cells. To confirm that Cap1₁₃₉₋₁₄₇ is presented on the surface of *Chlamydia* infected cells, Balb-3T3 (H-2^d) cells were infected with *C. trachomatis* serovar L2 and tested to determine whether these cells are recognized by a CD8+ T-cell clone specific for Cap1#139-147 epitope (SEQ ID NO: 145). The T-cell clone specific for Cap1#139-147 epitope was obtained by limiting dilution of the line 69 T-cells. The T-cell clone specifically recognized the *Chlamydia* infected cells. In these experiments, target cells were *C. trachomatis* infected (positive control) or uninfected Balb/3T3 cells, showing 45%, 36% and 30% specific lysis at 30:1, 10:1 and 3:1 effector to target ratios, respectively; or Cap1#139-147 epitope (SEQ ID NO: 145) coated, or untreated P815 cells, showing 83%, 75% and 58% specific lysis at 30:1, 10:1 and 3:1 effector to target ratios, respectively (negative controls having less than 5% lysis in all cases). This data suggests that the epitope is presented during infection.

109

In vivo studies show Cap1#139-147 epitope-specific T-cells are primed during murine infection with C. trachomatis. To determine if infection with C. trachomatis primes a Cap1#139-147 epitope-specific T-cell response, mice were infected i.p. with 108 IFU of C. trachomatis serovar L2. Two weeks after infection, the mice were sacrificed and spleen cells were stimulated on irradiated syngeneic spleen cells pulsed with Cap1#139-147 epitope peptide. After 5 days of stimulation, the cultures were used in a standard 51Cr release assay to determine if there were Cap1#139-147 epitope-specific T-cells present in the culture. Specifically, spleen cells from a C. trachomatis serovar L2 immunized mouse or a control mouse injected with PBS after a 5 days culture with Cap1#139-147 peptide-coated syngeneic spleen cells and CD8+ Tcells able to specifically recognize Cap1#139-147 epitope gave 73%, 60% and 32% specific lysis at a30:1, 10:1 and 3:1 effector to target ratios, respectively. The control mice had a percent lysis of approximately 10% at a 30:1 effector to target ratio, and steadily declining with lowering E:T ratios. Target cells were Cap1#139-147 peptidecoated, or untreated P815 cells. These data suggest that Cap1#139-147 peptide-specific T-cells are primed during murine infection with C. trachomatis.

Ct529 Localization

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Studies were performed demonstrating that Ct529 (referred to herein as Cap-1) localizes to the inclusion membrane of C. trachomatis-infected cells and is not 20 associated with elementary bodies or reticulate bodies. As described above, Cap-1 was identified as a product from Chlamydia that stimulates CD8+ CTL. These CTL are protective in a murine model of infection, thus making Cap-1 a good vaccine candidate. Further, since these CTL are MHC-I restricted, the Cap-1 gene must have access to the cytosol of infected cells, which may be a unique characteristic of specific Chlamydial 25 Therefore, determination of the cellular localization of the gene gene products. products would be useful in characterizing Cap-1 as a vaccine candidate. To detect the intracellular localization of Cap-1, rabbit polyclonal antibodies directed against a recombinant polypeptide encompassing the N-terminal 125 amino acids of Cap-1 (SEQ ID NO: 305, with the amino acid sequence including the N-terminal 6-His tag provided 30 in SEQ ID NO: 304) were used to stain McCoy cells infected with Chlamydiae.

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Rabbit-anti-Cap-1 polyclonal antibodies were obtained by hyper-immunization of rabbits with a recombinant polypeptide, rCt529c1-125 (SEQ ID NO: 305) encompassing the N-terminal portion of Cap-1. Recombinant rCt529e1-125 protein was obtained from *E. coli* transformed with a pET expression plasmid (as described above) encoding the nucleotides 1-375 encoding the N-terminal 1-125 amino acids of Cap-1. Recombinant protein was purified by Ni-NTA using techniques well known in the art. For a positive control antiserum, polyclonal antisera directed against elementary bodies were made by immunization of rabbits with purified *C. trachomatis* elementary bodies (Biodesign, Sacco, Maine). Pre-immune sera derived from rabbits prior to immunization with the Cap-1 polypeptide was used as a negative control.

Immunocytochemistry was performed on McCoy cell monolayers grown on glass coverslips inoculated with either *C. trachomatis* serovar L2 or *C. psitacci*, strain 6BC, at a concentration of 10⁶ IFU (Inclusion Forming Units) per ml. After 2 hours, medium was aspirated and replaced with fresh RP-10 medium supplemented with cycloheximide (1.0 μg/ml). Infected cells were incubated at in 7% CO₂ for 24 hours and fixed by aspirating medium, rinsing cells once with PBS and methanol fixation for 5 minutes. For antigen staining, fixed cell monolayers were washed with PBS and incubated at 37°C for 2 hours with 1:100 dilutions of specific or control antisera. Cells were rinsed with PBS and incubated for 1 hour with fluorescein isothiocyanate (FITC)-labeled, anti-rabbit IgG (KPL, Gaithersburg) and stained with Evans blue (0.05%) in PBS. Fluorescence was observed with a 100X objective (Zeiss epifluorescence microscope), and photographed (Nikon UFX-11A camera).

Results from this study show Cap-1 localizes to the inclusion membrane of *C. trachomatis*-infected cells. Cap-1 specific antibody labeled the inclusion membranes of *C. trachomatis*-infected cells, but not *Chlamydial* elementary bodies contained in these inclusions or released by the fixation process. Conversely, the anti-elementary body antibody clearly labeled the bacterial bodies, not only within the inclusions, but those released by the fixation process. Specificity of the anti-Cap-1 antibody is demonstrated by the fact that it does not stain *C. psittaci*-infected cells. Specificity of the Cap-1 labeling is also shown by the absence of reactivity in pre-immune sera. These results suggest that Cap-1 is released from the bacteria and becomes associated with the *Chlamydial* inclusion membrane. Therefore, Cap-1 is a

111

gene product which may be useful for stimulating CD8+ T cells in the development of a vaccine against infections caused by *Chlamydia*.

The relevance of the Cap-1 gene as a potential CTL antigen in a vaccine against *Chlamydia* infection is further illustrated by two additional series of studies. First, CTL specific for the MHC-I epitope of Cap-1 CT529 #138-147 peptide of *C. trachomatis* (SEQ ID NO: 144) have been shown to be primed to a high frequency during natural infection. Specifically, Balb/C mice were inoculated with 10⁶ I.F.U. of *C. trachomatis*, serova L2. After 2 weeks, spleens were harvested and quantified by Elispot analysis for the number of IFN-γ secreting cells in response to Cap-1 #138-147 peptide-pulsed antigen presenting cells. In two experiments, the number of IFN-γ-secreting cells in 10⁵ splenocytes was about 1% of all CD8+ T-cells. This high frequency of responding CD8+ CTL to the MHC-1 epitope (Cap-1 CT529 #138-147 peptide) suggest that Cap-1 is highly immunogenic in infections.

Results from a second series of studies have shown that the Cap-1 protein is almost immediately accessible to the cytosol of the host cell upon infection. This is shown in a time-course of Cap-1 CT529 #138-147 peptide presentation. Briefly, 3T3 cells were infected with *C. trachomatis* serovar L2 for various lengths of time, and then tested for recognition by Cap-1 CT529 #138-147 peptide-specific CTL. The results show that *C. trachomatis*-infected 3T3 cells are targeted for recognition by the antigen-specific CTL after only 2 hours of infection. These results suggest that Cap-1 is an early protein synthesized in the development of *C. trachomatis* elementary bodies to reticulate bodies. A CD8+ CTL immune response directed against a gene product expressed early in infection may be particularly efficacious in a vaccine against *Chlamydia* infection.

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EXAMPLE 5

GENERATION OF ANTIBODY AND T-CELL RESPONSES IN MICE IMMUNIZED WITH CHLAMYDIA ANTIGENS

Immunogenicity studies were conducted to determine the antibody and CD4+ T cell responses in mice immunized with either purified SWIB or S13 proteins formulated with Montanide adjuvant, or DNA-based immunizations with pcDNA-3 expression

112

vectors containing the DNA sequences for SWIB or S13. SWIB is also referred to as clone 1-B1-66 (SEQ ID NO: 1, with the corresponding amino acid sequence provided in SEQ ID NO: 5), and S13 ribosomal protein is also referred to as clone 10-C10-31 (SEQ ID NO: 4, with the corresponding amino acid sequence provided in SEO ID NO: 12). In the first experiment, groups of three C57BL/6 mice were immunized twice and monitored for antibody and CD4+ T-cell responses. DNA immunizations were intradermal at the base of the tail and polypeptide immunizations were administered by subcutaneous route. Results from standard ³H-incorporation assays of spleen cells from immunized mice shows a strong proliferative response from the group immunized with purified recombinant SWIB polypeptide (SEQ ID NO: 5). Further analysis by cytokine induction assays, as previously described, demonstrated that the group immunized with SWIB polypeptide produced a measurable IFN-y and IL-4 response. Subsequent ELISA-based assays to determine the predominant antibody isotype response in the experimental group immunized with the SWIB polypeptide were performed. Fig. 4 illustrates the SWIB-immunized group gave a humoral response that was predominantly IgG1.

In a second experiment, C3H mice were immunized three times with 10 μg purified SWIB protein (also referred to as clone 1-B1-66, SEQ ID NO: 5) formulated in either PBS or Montanide at three week intervals and harvested two weeks after the third immunization. Antibody titers directed against the SWIB protein were determined by standard ELISA-based techniques well known in the art, demonstrating the SWIB protein formulated with Montanide adjuvant induced a strong humoral immune response. T-cell proliferative responses were determined by a XTT-based assay (Scudiero, et al, *Cancer Research*, 1988, 48:4827). As shown in Fig. 5, splenocytes from mice immunized with the SWIB polypeptide plus Montanide elicited an antigen specific proliferative response. In addition, the capacity of splenocytes from immunized animals to secrete IFN-γ in response to soluble recombinant SWIB polypeptide was determined using the cytokine induction assay previously described. The splenocytes from all animals in the group immunized with SWIB polypeptide formulated with montanide adjuvant secreted IFN-γ in response to exposure to the SWIB Chlamydia antigen, demonstrating an *Chlamydia*-specific immune response.

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In a further experiment, C3H mice were immunized at three separate time points at the base of the tail with 10 μg of purified SWIB or S13 protein (*C. trachomatis*, SWIB protein, clone 1-B1-66, SEQ ID NO: 5, and S13 protein, clone 10-C10-31, SEQ ID NO: 4) formulated with the SBAS2 adjuvant (SmithKline Beecham, London, England). Antigen-specific antibody titers were measured by ELISA, showing both polypeptides induced a strong IgG response, ranging in titers from 1 x10⁻⁴ to 1 x10⁻⁵. The IgG1 and IgG2a components of this response were present in fairly equal amounts. Antigen-specific T-cell proliferative responses, determined by standard ³H-incorporation assays on spleen cells isolated from immunized mice, were quite strong for SWIB (50,000 cpm above the negative control) and even stronger for s13 (100,000 cpm above the negative control). The IFNγ production was assayed by standard ELISA techniques from supernatant from the proliferating culture. *In vitro* restimulation of the culture with S13 protein induced high levels of IFNγ production, approximately 25 ng/ml versus 2 ng/ml for the negative control. Restimulation with the SWIB protein also induced IFNγ, although to a lesser extent.

In a related experiment, C3H mice were immunized at three separate time points with 10 μg of purified SWIB or S13 protein (*C. trachomatis*, SWIB protein, clone 1-B1-66, SEQ ID NO: 5, and S13 protein, clone 10-C10-31, SEQ ID NO: 4) mixed with 10 μg of Cholera Toxin. Mucosal immunization was through intranasal inoculation. Antigen-specific antibody responses were determined by standard ELISA techniques. Antigen-specific IgG antibodies were present in the blood of SWIB-immunized mice, with titers ranging from 1 x10⁻³ to 1 x10⁻⁴, but non-detectable in the S13-immunized animals. Antigen-specific T-cell responses from isolated splenocytes, as measured by IFNγ production, gave similar results to those described immediately above for systemic immunization.

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An animal study was conducted to determine the immunogenicity of the CT529 serovar LGVII CTL epitope, defined by the CT529 10mer consensus peptide (CSFIGGITYL – SEQ ID NO: 31), which was identified as an H2-Kd restricted CTL epitope. BALB/c mice (3 mice per group) were immunized three times with 25 µg of peptide combined with various adjuvants. The peptide was administered systemically at the base of the tail in either SKB Adjuvant System SBAS-2", SBAS-7 (SmithKline

Beecham, London, England) or Montanide. The peptide was also administered intranasally mixed with 10ug of Cholera Toxin (CT). Naive mice were used as a control. Four weeks after the 3rd immunization, spleen cells were restimulated with LPS-blasts pulsed with 10ug/ml CT529 10mer consensus peptide at three different effector to LPS-blasts ratios: 6, 1.5 and 0.4 at 1x10⁶ cell/ml. After 2 restimulations, effector cells were tested for their ability to lyse peptide pulsed P815 cells using a standard chromium release assay. A non-relevant peptide from chicken egg ovalbumin was used as a negative control. The results demonstrate that a significant immune response was elicited towards the CT529 10mer consensus peptide and that antigenspecific T-cells capable of lysing peptide-pulsed targets were elicited in response to immunization with the peptide. Specifically, antigen-specific lytic activities were found in the SBAS-7 and CT adjuvanted group while Montanide and SBAS-2" failed to adjuvant the CTL epitope immunization.

EXAMPLE 6

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EXPRESSION AND CHARACTERIZATION OF CHLAMYDIA PNEUMONIAE GENES

The human T-cell line, TCL-8, described in Example 1, recognizes Chlamydia trachomatis as well as Chlamydia pneumonia infected monocyte-derived dendritic cells, suggesting Chlamydia trachomatis and pneumonia may encode cross-reactive T-cell epitopes. To isolate the Chlamydia pneumonia genes homologous to Chlamydia trachomatis LGV II clones 1B1-66, also referred to as SWIB (SEQ ID NO: 1) and clone 10C10-31, also referred to as S13 ribosomal protein (SEQ ID NO: 4), HeLa 229 cells were infected with C. pneumonia strain TWAR (CDC/CWL-029). After three days incubation, the C. pneumonia-infected HeLa cells were harvested, washed and resuspended in 200 µl water and heated in a boiling water bath for 20 minutes. Ten microliters of the disrupted cell suspension was used as the PCR template.

C. pneumonia specific primers were designed for clones 1B1-66 and 10C10-31 such that the 5' end had a 6X-Histidine tag and a Nde I site inserted, and the 3' end had a stop codon and a BamHI site included (Fig. 6). The PCR products were amplified and sequenced by standard techniques well known in the art. The C.

115

pneumonia-specific PCR products were cloned into expression vector pET17B (Novagen, Madison, WI) and transfected into E. coli BL21 pLysS for expression and subsequent purification utilizing the histidine-nickel chromatographic methodology provided by Novagen. Two proteins from *C. pneumonia* were thus generated, a 10-11 kDa protein referred to as CpSWIB (SEQ ID NO: 27, and SEQ ID NO: 78 having a 6X His tag, with the corresponding amino acid sequence provided in SEQ ID NO: 28, respectively), a 15 kDa protein referred to as CpS13 (SEQ ID NO: 29, and SEQ ID NO: 77, having a 6X His tag, with the corresponding amino acid sequence provided in SEQ ID NO: 30 and 91, respectively).

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EXAMPLE 7

INDUCTION OF T CELL PROLIFERATION AND INTERFERON-γ PRODUCTION BY CHLAMYDIA PNEUMONIAE ANTIGENS

The ability of recombinant *Chlamydia pneumoniae* antigens to induce T cell proliferation and interferon-γ production is determined as follows.

Proteins are induced by IPTG and purified by Ni-NTA agarose affinity chromatography (Webb et al., *J. Immunology 157*:5034-5041, 1996). The purified polypeptides are then screened for the ability to induce T-cell proliferation in PBMC preparations. PBMCs from *C. pneumoniae* patients as well as from normal donors whose T-cells are known to proliferate in response to *Chlamydia* antigens, are cultured in medium comprising RPMI 1640 supplemented with 10% pooled human serum and 50 μg/ml gentamicin. Purified polypeptides are added in duplicate at concentrations of 0.5 to 10 μg/mL. After six days of culture in 96-well round-bottom plates in a volume of 200 μl, 50 μl of medium is removed from each well for determination of IFN-γ levels, as described below. The plates are then pulsed with 1 μCi/well of tritiated thymidine for a further 18 hours, harvested and tritium uptake determined using a gas scintillation counter. Fractions that result in proliferation in both replicates three fold greater than the proliferation observed in cells cultured in medium alone are considered positive.

IFN-y was measured using an enzyme-linked immunosorbent assay (ELISA). ELISA plates are coated with a mouse monoclonal antibody directed to human IFN-y (PharMingen, San Diego, CA) in PBS for four hours at room temperature. Wells are then blocked with PBS containing 5% (W/V) non-fat dried milk for 1 hour at room temperature. The plates are washed six times in PBS/0.2% TWEEN-20 and samples diluted 1:2 in culture medium in the ELISA plates are incubated overnight at room temperature. The plates are again washed and a polyclonal rabbit anti-human IFN-y serum diluted 1:3000 in PBS/10% normal goat serum is added to each well. The plates are then incubated for two hours at room temperature, washed and horseradish peroxidase-coupled anti-rabbit IgG (Sigma Chemical So., St. Louis, MO) is added at a 1:2000 dilution in PBS/5% non-fat dried milk. After a further two hour incubation at room temperature, the plates are washed and TMB substrate added. The reaction is stopped after 20 min with 1 N sulfuric acid. Optical density is determined at 450 nm using 570 nm as a reference wavelength. Fractions that result in both replicates giving an OD two fold greater than the mean OD from cells cultured in medium alone, plus 3 standard deviations, are considered positive.

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A human anti-Chlamydia T-cell line (TCL-8) capable of cross-reacting to C. trachomatis and C. pneumonia was used to determine whether the expressed proteins described in the example above, (i.e., CpSWIB, SEQ ID NO: 27, and SEQ ID NO: 78 having a 6X His tag, with the corresponding amino acid sequence provided in SEQ ID NO: 28, respectively, and the 15 kDa protein referred to as CpS13 SEQ ID NO: 29, and SEQ ID NO: 77, having a 6X His tag, with the corresponding amino acid sequence provided in SEQ ID NO: 30 and 91, respectively), possessed T-cell epitopes common to both C. trachomatis and C. pneumonia. Briefly, E. coli expressing Chlamydial proteins were titered on 1 x 10⁴ monocyte-derived dendritic cells. After two hours, the dendritic cells cultures were washed and 2.5 x 10⁴ T cells (TCL-8) added and allowed to incubate for an additional 72 hours. The amount of INF-γ in the culture supernatant was then determined by ELISA. As shown in Figs. 7A and 7B, the TCL-8 T-cell line specifically recognized the S13 ribosomal protein from both C. trachomatis and C. pneumonia as demonstrated by the antigen-specific induction of IFN-γ, whereas only the SWIB protein from C. trachomatis was recognized by the T-cell line. To

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validate these results, the T cell epitope of *C. trachomatis* SWIB was identified by epitope mapping using target cells pulsed with a series of overlapping peptides and the T-cell line TCL-8. 3H-thymidine incorporation assays demonstrated that the peptide, referred to as C.t.SWIB 52-67, of SEQ ID NO: 39 gave the strongest proliferation of the TCL-8 line. The homologous peptides corresponding to the SWIB of *C. pneumoniae* sequence (SEQ ID NO: 40), the topoisomerase-SWIB fusion of *C. pneumoniae* (SEQ ID NO: 43) and *C. trachomatis* (SEQ ID NO: 42) as well as the human SWI domain (SEQ ID NO: 41) were synthesized and tested in the above assay. The T-cell line TCL-8 only recognized the *C. trachomatis* peptide of SEQ ID NO: 39 and not the corresponding *C. pneumoniae* peptide (SEQ ID NO: 40), or the other corresponding peptides described above (SEQ ID NO: 41-43).

117

Chlamydia-specific T cell lines were generated from donor CP-21 with a positive serum titer against *C. pneumoniae* by stimulating donor PBMC with either *C. trachomatis* or *C. pneumoniae*-infected monocyte-derived dendritic cells, respectively.

T-cells generated against *C. pneumoniae* responded to recombinant *C. pneumoniae*-SWIB but not *C. trachomatis*-SWIB, whereas the T-cell line generated against *C. trachomatis* did not respond to either *C. trachomatis*- or *C. pneumoniae*-SWIB (see Fig. 9). The *C. pneumoniae*-SWIB specific immune response of donor CP-21 confirms the *C. pneumoniae* infection and indicates the elicitation of *C. pneumoniae*-SWIB specific T-cells during *in vivo C. pneumoniae* infection.

Epitope mapping of the T-cell response to *C. pneumoniae*-SWIB has shown that Cp-SWIB-specific T-cells responded to the overlapping peptides Cp-SWIB 32-51 (SEQ ID NO: 101) and Cp-SWIB 37-56 (SEQ ID NO: 102), indicating a *C. pneumoniae*-SWIB-specific T-cell epitope Cp-SWIB 37-51 (SEQ ID NO: 100).

In additional experiments, T-cell lines were generated from donor CP1, also a *C. pneumoniae* seropositive donor, by stimulating PBMC with non-infectious elementary bodies from *C. trachomatis* and *C. pneumoniae*, respectively. In particular, proliferative responses were determined by stimulating 2.5 x 10⁴ T-cells in the presence of 1 x 10⁴ monocyte-derived dendritic cells and non-infectious elementary bodies derived from *C. trachomatis* and *C. pneumoniae*, or either recombinant *C. trachomatis* or *C. pneumoniae* SWIB protein. The T-cell response against SWIB resembled the data obtained with T-cell lines from CP-21 in that *C. pneumoniae*-SWIB, but not *C.*

118

trachomatis-SWIB elicited a response by the C. pneumoniae T-cell line. In addition, the C. trachomatis T-cell line did not proliferate in response to either C. trachomatis or C. pneumoniae SWIB, though it did proliferate in response to both CT and CP elementary bodies. As described in Example 1, Clone 11-C12-91 (SEQ ID NO: 63), identified using the TCP-21 cell line, has a 269 bp insert that is part of the OMP2 gene (CT443) and shares homology with the 60 kDa cysteine rich outer membrane protein of C. pneumoniae, referred to as OMCB. To further define the reactive epitope(s), epitope mapping was performed using a series of overlapping peptides and the immunoassay previously described. Briefly, proliferative responses were determined by stimulating 2.5 x 10⁴ TCP-21 T-cells in the presence of 1 x 10⁴ monocyte-derived dendritic cells with either non-infectious elementary bodies derived from C. trachomatis and C. pneumoniae, or peptides derived from the protein sequence of C. trachomatis or C. pneumoniae OMCB protein (0.1 µg/ml). The TCP-21 T-cells responded to epitopes CT-OMCB #167-186, CT-OMCB #171-190, CT-OMCB #171-186, and to a lesser extent, CT-OMCB #175-186 (SEQ ID NO: 249-252, respectively). Notably, the TCP-21 T-cell line also gave a proliferative response to the homologous C. pneumoniae peptide CP-OMCB #171-186 (SEQ ID NO: 253), which was equal to or greater than the response to the to the C. trachomatis peptides. The amino acid substitutions in position two (i.e., Asp for Glu) and position four (i.e., Cys for Ser) did not alter the proliferative response of the T-cells and therefore demonstrating this epitope to be a cross-reactive epitope between C. trachomatis and C. pneumoniae.

EXAMPLE 8 IMMUNE RESPONSES OF HUMAN PBMC AND T-CELL LINES AGAINST CHLAMYDIA ANTIGENS

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The examples provided herein suggest that there is a population of healthy donors among the general population that have been infected with *C. trachomatis* and generated a protective immune response controlling the *C. trachomatis* infection. These donors remained clinically asymptomatic and seronegative for *C. trachomatis*. To characterize the immune responses of normal donors against chlamydial antigens which had been identified by CD4 expression cloning, PBMC

119

obtained from 12 healthy donors were tested against a panel of recombinant chlamydial antigens including C. trachomatis-, C. pneumoniae-SWIB and C. trachomatis-, C. pneumoniae-S13. The data are summarized in Table I below. All donors were seronegative for C. trachomatis, whereas 6/12 had a positive C. pneumoniae titer. Using a stimulation index of >4 as a positive response, 11/12 of the subjects responded to C. trachomatis elementary bodies and 12/12 responded to C. pneumoniae elementary bodies. One donor, AD104, responded to recombinant C. pneumoniae-S13 protein, but not to recombinant C. trachomatis-S13 protein, indicating a C. pneumoniae-specific response. Three out of 12 donors had a C. trachomatis-SWIB, but not a C. pneumoniae-SWIB specific response, confirming a C. trachomatis infection. C. trachomatis and C. pneumoniae-S13 elicited a response in 8/12 donors suggesting a chlamydial infection. These data demonstrate the ability of SWIB and S13 to elicit a T-cell response in PBMC of normal study subjects.

120

Table I.

Immune response of normal study subjects against Chlamydia

Donar	Sex	<i>Chlanydia</i> IgGtiter	CT EB	CP EB	CT Swib	CP Swib	CT S13	CP SI3	CT lpdA	CT TSA
4D100	male	negative	++	+++	+	-	++	++	_	nt.
4D104	female	negative	+++	++-	-	_	-	++	-	nt.
4D108	male	CP 1:256	++	++	+	+/-	+	+	+	nt.
4D112	female	negative	++	++	+	-	+	-	+/-	nt.
AD120	male	negative	-	+	-	-	-	-	-	nt
4 D124	female	CP 1:128	++	++	-	-	-	-	-	nt.
4D128	male	CP 1:512	+	++	-	-	++	+	++	-
4D132	female	negative	++	++	-	-	+	+	-	-
4D136	female	CP 1:128	+	+	-	-	+/-	-	-	-
4D140	male	CP 1:256	++	++	-	-	+	+	-	-
4D142	female	CP 1:512	++	++	-	-	+	+	+	-
4D146	female	negative	+1-	++	-		++	+	+	

CT= Chlamydia trachomatis; CP= Chlamydia pneumoniae; EB= Chlamydia elementary bodies; Swib= recombinant Chlamydia Swib protein; S13= recombinant Chlamydia S13 protein; lpdA= recombinant Chlamydia lpdA protein; TSA= recombinant Chlamydia TSA protein. Values represent results from standard proliferation assays. Proliferative responses were determined by stimulating 3 x 10⁵ PBMC with 1 x 10⁴ monocyte-derived dendritic cells pre-incubated with the respective recombinant antigens or elementary bodies (EB). Assays were harvested after 6 days with a ³H-thymidine pulse for the last 18h.

SI: Stimulation index

+/-: SI ~ 4
15 +: SI > 4
++: SI 10-30
+++: SI > 30

121

In a first series of experiments, T-cell lines were generated from a healthy female individual (CT-10) with a history of genital exposure to *C. trachomatis* by stimulating T-cells with *C. trachomatis* LGV II elementary bodies as previously described. Although the study subject was exposed to *C. trachomatis*, she did not seroconvert and did not develop clinical symptoms, suggesting donor CT-10 may have developed a protective immune response against *C. trachomatis*. As shown in Fig. 10, a primary *Chlamydia*-specific T-cell line derived from donor CT-10 responded to *C. trachomatis*-SWIB, but not *C. pneumoniae*-SWIB recombinant proteins, confirming the exposure of CT-10 to *C. trachomatis*. Epitope mapping of the T-cell response to *C. trachomatis*-SWIB showed that this donor responded to the same epitope Ct-SWIB 52-67 (SEQ ID NO: 39) as T-cell line TCL-8, as shown in Fig. 11.

Additional T-cell lines were generated as described above for various *C. trachomatis* patients. A summary of the patients' clinical profile and proliferative responses to various *C. trachomatis* and *C. pneumoniae* elementary bodies and recombinant proteins are summarized in Table II as follows:

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	Prolifer	ative respo	nse of	C. tre	achom	atis pa	tient	<u> </u>		
?atients	Clinical manifestation	IgG titer	CT EB	CP EB	CT Swib	CP Swib	CT S13	CP S13	CT lpdA	CT TSA
CT-1	NGU	negative	+	+	-	-	++	++	++	+
CT-2	NGU	negative	++	++	-	_	+	+/-	-	-
CT-3	asymptomatic shed Eb Dx was HPV	Ct 1:512 Cp 1:1024 Cps 1:256	+	+	-	-	+	-	+	-
СТ-4	asymptomatic shed Eb	Ct 1:1024	+	+	-	-	-	-	-	-
CT-5	BV	Ct 1:256 Cp 1:256	++	++	-	-	+	-	-	-
СТ-6	perinial rash discharge	Cp 1:1024	+	+	-	-	-	-	-	-
CT-7	BV genital ulcer	Ct 1:512 Cp 1:1024	+	+	-	-	+	+	+	-
CT-8	Not known	Not tested	++	++	- -	-	-	•	-	-
CT-9	asymptomatic	Ct 1:128 Cp 1:128	+++	++	-	-	++	+	+	-
СТ-10	Itch mild vulvar	negative	++	++	-	-	-	-	-	-
CT-11	BV, abnormal pap	Ct 1: 512	+++	+++	-	-	+++	+/-	++	+
CT-12	asymptomatic	Cp 1: 512	++	++	_	-	++	+	+	_

NGU= Non-Gonococcal Urethritis; BV= Bacterial Vaginosis; CT= Chlamydia trachomatis; CP= Chlamydia pneumoniae; EB= Chlamydia elementary bodies; Swib= recombinant Chlamydia Swib protein; S13= recombinant Chlamydia S13 protein; lpdA= recombinant Chlamydia lpdA protein; TSA= recombinant Chlamydia TSA protein

Values represent results from standard proliferation assays. Proliferative responses were determined by stimulating 3×10^5 PBMC with the respective recombinant antigens or elementary bodies (EB). Assays were harvested after 6 days with a 3 H-thymidine pulse for the last 18 hours.

SI: Stimulation index

123

Using the panel of asymptomatic (as defined above) study subjects and *C. trachomatis* patients, as summarized in Tables I and II, a comprehensive study of the immune responses of PBMC derived from the two groups was conducted. Briefly, PBMCs from *C. pneumoniae* patients as well as from normal donors are cultured in medium comprising RPMI 1640 supplemented with 10% pooled human serum and 50 μ g/ml gentamicin. Purified polypeptides, a panel of recombinant *chlamydial* antigens including *C. trachomatis-, C. pneumoniae-*SWIB and S13, as well as . *C. trachomatis* lpdA and TSA are added in duplicate at concentrations of 0.5 to 10 μg/mL. After six days of culture in 96-well round-bottom plates in a volume of 200 μl, 50 μl of medium is removed from each well for determination of IFN-γ levels, as described below. The plates are then pulsed with 1 μCi/well of tritiated thymidine for a further 18 hours, harvested and tritium uptake determined using a gas scintillation counter. Fractions that result in proliferation in both replicates three fold greater than the proliferation observed in cells cultured in medium alone are considered positive.

Proliferative responses to the recombinant *Chlamydiae* antigens demonstrated that the majority of asymptomatic donors and *C. trachomatis* patients recognized the *C. trachomatis* S13 antigen (8/12) and a majority of the *C. trachomatis* patients recognized the *C. pneumonia* S13 antigen (8/12), with 4/12 asymptomatic donors also recognizing the *C. pneumonia* S13 antigen. Also, six out of twelve of the *C. trachomatis* patients and four out of twelve of the asymptomatic donors gave a proliferative response to the lpdA antigen of *C. trachomatis*. These results demonstrate that the *C. trachomatis* and *C. pneumonia* S13 antigen, *C. trachomatis* Swib antigen and the *C. trachomatis* lpdA antigen are recognized by the asymptomatic donors, indicating these antigens were recognized during exposure to *Chlamydia* and an immune response elicited against them. This implies these antigens may play a role in conferring protective immunity in a human host. In addition, the *C. trachomatis* and *C. pneumonia* S13 antigen is recognized equally well among the *C. trachomatis* patients, therefore indicating there may be epitopes shared between *C. trachomatis* and *C. pneumonia* in the S13 protein. Table III summarizes the results of these studies.

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Table III.

Antigen	Normal Donors	C.t. Patients
C.tSwib	3/12	0/12
C.pSwib	0/12	0/12
C.tS13	8/12	8/12
C.pS13	4/12	8/12
lpdA	4/12	6/12
TSA	0/12	2/12

A series of studies were initiated to determine the cellular immune response to short-term T-cell lines generated from .asymptomatic donors and C. trachomatis patients. Cellular immune responses were measured by standard proliferation assays and IFN-γ, as described in Example 7. Specifically, the majority of the antigens were in the form of single E. coli clones expressing Chlamydial antigens, although some recombinant proteins were also used in the assays. The single E. coli clones were titered on 1 x 104 monocyte-derived dendritic cells and after two hours, the culture was washed and 2.5 x 10⁴ T-cells were added. The assay using the recombinant proteins were performed as previously described. Proliferation was determined after four days with a standard ³H-thymidine pulse for the last 18 hours. Induction of IFN-y was determined from culture supernatants harvested after four days using standard ELISA assays, as described above. The results show that all the C. trachomatis antigens tested, except for C.T. Swib, elicited a proliferative response from one or more different T-cell lines derived form C. trachomatis patients. In addition, proliferative responses were elicited from both the C. trachomatis patients and asymptomatic donors for the following Chlamydia genes, CT622, groEL, pmpD, CT610 and rS13.

The 12G3-83 clone also contains sequences to CT734 and CT764 in addition to CT622, and therefore these gene sequence may also have immunoreactive epitopes. Similarly, clone 21G12-60 contains sequences to the hypothetical protein genes CT229 and CT228 in addition to CT875; and 15H2-76 also contains sequences

from CT812 and CT088, as well as sharing homology to the sycE gene. Clone 11H3-61 also contains sequences sharing homology to the PGP6-D virulence protein.

Table IV.

Clone	C. t.	TCL from	TCL from	SEQ ID NO:
	Antigen	Asymp. Donors	C. t.	
	(putative*)		Patients	
1B1-66 (E. coli)	Swib	2/2	0/4	5
1B1-66 (protein)	Swib	2/2	0/4	5
12G3-83 (E. coli)	CT622*	2/2	4/4	57
22B3-53 (E. coli)	groEL	1/2	4/4	111
22B3-53 (protein)	groEL	1/2	4/4	111
15H2-76 (E. coli)	PmpD*	1/2	3/4	87
11H3-61 (E. coli)	rL1*	0/2	3/4	60
14H1-4 (E. coli)	TSA	0/2	3/4	56
14H1-4 (protein)	TSA	0/2	3/4	56
11G10-46 (E. coli)	CT610	1/2	1/4	62
10C10-17 (E. coli)	rS13	1/2	1/4	62
10C10-17 (protein)	rS13	1/2	1/4	62
21G12-60 (E. coli)	CT875*	0/2	2/4	110
11H4-32 (E. coli)	dnaK	0/2	2/4	59
21C7-8 (E. coli)	dnaK	0/2	2/4	115
17C10-31 (E. coli)	CT858	0/2	2/4	114

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EXAMPLE 9 PROTECTION STUDIES USING CHLAMYDIA ANTIGENS

1. SWIB

Protection studies were conducted in mice to determine whether immunization with chlamydial antigens can impact on the genital tract disease resulting from chlamydial inoculation. Two models were utilized; a model of intravaginal inoculation

126

that uses a human isolate containing a strain of Chlamydia psittaci (MTW447), and a model of intrauterine inoculation that involves a human isolate identified as Chlamydia trachomatis, serovar F (strain NI1). Both strains induce inflammation in the upper genital tract, which resemble endometritis and salpingitis caused by Chlamvdia trachomatis in women. In the first experiment, C3H mice (4 mice per group) were immunized three times with 100 µg of pcDNA-3 expression vector containing C. trachomatis SWIB DNA (SEQ ID NO: 1, with the corresponding amino acid sequence provided in SEQ ID NO: 5). Inoculations were at the base of the tail for systemic immunization. Two weeks after the last immunization, animals were progesterone treated and infected, either thru the vagina or by injection of the inoculum in the uterus. Two weeks after infection, the mice were sacrificed and genital tracts sectioned, stained and examined for histopathology. Inflammation level was scored (from + for very mild, to +++++ for very severe). Scores attributed to each single oviduct /ovary were summed and divided by the number of organs examined to get a mean score of inflammation for the group. In the model of uterine inoculation, negative controlimmunized animals receiving empty vector showed consistent inflammation with an ovary /oviduct mean inflammation score of 6.12, in contrast to 2.62 for the DNAimmunized group. In the model of vaginal inoculation and ascending infection, negative control-immunized mice had an ovary /oviduct mean inflammation score of 8.37, versus 5.00 for the DNA-immunized group. Also, in the later model, vaccinated mice showed no signs of tubal occlusion while negative control vaccinated groups had inflammatory cells in the lumen of the oviduct

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In a second experiment, C3H mice (4 mice per group) were immunized three times with 50 µg of pcDNA-3 expression vector containing *C. trachomatis* SWIB DNA (SEQ ID NO: 1, with the corresponding amino acid sequence provided in SEQ ID NO: 5) encapsulated in Poly Lactide co-Glycolide microspheres (PLG); immunizations were made intra-peritoneally. Two weeks after the last immunization, animal were progesterone treated and infected by inoculation of *C. psittaci* in the vagina. Two weeks after infection, mice were sacrificed and genital tracts sectioned, stained and examined for histopathology. Inflammation level was scored as previously described. Scores attributed to each single oviduct /ovary were summed and divided by the number of examined organs to get a mean of inflammation for the group. Negative control-

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immunized animals receiving PLG-encapsulated empty vector showed consistent infammation with an ovary /oviduct mean inflammation score of 7.28, versus 5.71 for the PLG-encapsulated DNA immunized group. Inflammation in the peritoneum was 1.75 for the vaccinated group versus 3.75 for the control.

127

In a third experiment, C3H mice (4 per group) were immunized three times with 10 µg of purified recombinant protein, either SWIB (SEQ ID NO: 1, with the corresponding amino acid sequence provided in SEQ ID NO: 5, or S13 (SEQ ID NO: 4, with the corresponding amino acid sequence provided in SEQ ID NO: 12) mixed with Cholera Toxin (CT); the preparation was administred intranasally upon anaesthesia in a 20 uL volume. Two weeks after the last immunization, animal were progesterone treated and infected, either by vaginal inoculation of C. psittaci or by injection of C. trachomatis serovar F in the uterus. Two weeks after infection, the mice were sacrificed and genital tracts sectioned, stained and examined for histopathology. The degree of inflammation was scored as described above. Scores attributed to each single oviduct /ovary were summed and divided by the number of examined organs to get a mean score of inflammation for the group. In the model of uterine inoculation, negative control- immunized animals receiving cholera toxin alone showed an ovary /oviduct mean inflammation score of 4.25 (only 2 mice analyzed; 2 other died) versus 5.00 for the s13 plus cholera toxin-immunized group, and 1.00 for the SWIB plus cholera toxin. Untreated infected animals had an ovary /oviduct mean inflammation score of 7. In the model of vaginal inoculation and ascending infection, negative control-immunized mice had an ovary/oviduct mean inflammation score of 7.37 versus 6.75 for the s13 plus cholera toxin-immunized group and 5.37 for the SWIB plus cholera toxin-immunized group. Untreated infected animals had an ovary /oviduct mean inflammation score of 8.

The three experiments described above suggest that SWIB-specific protection is obtainable. This protective effect is more marked in the model of homologous infection but is still present when in a heterologous challenge infection with *C. psittaci*.

PCT/US01/23121

2. CT529/Cap1

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CT529/Cap1 was identified earlier as a product from Chlamydia that stimulates CD8+ CTL. In this example, we sought to confirm that immunization with Cap1 would be protective in an animal model of chlamydia infection.

To generate recombinant vaccinia virus for delivery of a Capl immunogenic fragment, a DNA fragment containing a modified Kozak sequence and base pairs 319-530 of the cap1 gene (CT529) was amplified from C. trachomatis L2 genomic DNA using PCRTM and ligated into pSC11ss (Earl PL, Koenig S, Moss B (1991) Biological and immunological properties of human immunodeficiency virus type 1 envelope glycoprotein: analysis of proteins with truncations and deletions expressed by recombinant vaccinia viruses. J Virol. 65:31-41). DNA digested with SalI and StuI. The portion of the cap1 gene ligated into pSC11ss encodes amino acids 107-176 of Cap1 protein, containing the previously identified CTL epitope of amino acids 139-147. The resulting plasmid was used to transfect CV-1 cells (ATCC# CCL-70; Jensen FC et al. (1964) Infection of human and simian tissue cultures with Rous Sarcoma Virus. Proc. Natl. Acad. Sci. USA 52: 53-59.) which were subsequently infected with wildtype vaccinia virus. Homologous recombination between the wild-type virus and plasmid DNA generated recombinant vaccinia viruses which were selected on the basis of both beta-galactosidase expression and the inactivation of thymidine kinase, as described previously (Chakrabarti et al, Mol Cell Biol. 1985, 5(12):3403-9). Recombinant virus was plaque purified three times and titered after growth in human TK-143B cells. Virus preparations were treated with equal volume of 0.25 mg/ml trypsin for 30 mins. at 37°C and diluted in PBS prior to immunization of mice. Groups of 5 mice were used for all experimental and control groups. The data presented below are representative of three independent experiments.

A group of mice was immunized with 10⁶ of the recombinant vaccinia i.p. and was allowed to recover for 3 weeks. Negative control groups were immunized with either buffer alone or wild-type vaccinia. As a positive control, a group of mice was infected i.v. with 10⁶ i.f.u. of C. trachomatis. The number of organisms given to the positive control group has been previously shown to be cleared within 2 weeks. After 3 weeks, animals in each of the groups were challenged i.v. with 10⁶ i.f.u. of C.

trachomatis. Three days after challenge the mice were sacrificed and the number of i.f.u. per spleen was determined.

The mean number of organisms found in the spleens of animals immunized with the vaccinia virus expressing Cap1 (7.1x10⁴) was 2.6-fold fewer (p<0.01; Wilcoxon's-Rank Sum analysis) than animals in the control groups immunized with either buffer (1.8x10⁵) or wild-type vaccinia (1.9x10⁵). Animals in the positive group had 77-fold fewer organisms (2.4x103) per spleen than animals in the negative control groups (p<0.01; Wilcoxon's-Rank Sum analysis). These data demonstrate that immunization with an immunogenic fragment of Cap1 can afford a statistically significant level of protection against C. trachomatis infection.

EXAMPLE 10 Pmp/Ra12 FUSION PROTEINS

Various Pmp/Ra12 fusion constructs were generated by first synthesizing PCR fragments of a Pmp gene using primers containing a Not I restriction site. Each PCR fragment was then ligated into the NotI restriction site of pCRX1. The pCRX1 vector contains the 6HisRa12 portion of the fusion. The Ra12 portion of the fusion construct encodes a polypeptide corresponding to amino acid residues 192-323 of *Mycobacterium tuberculosis* MTB32A, as described in U.S. Patent Application 60/158,585, the disclosure of which is incorporated herein by reference. The correct orientation of each insert was determined by its restriction enzyme pattern and its sequence was verified. Multiple fusion constructs were made for PmpA, PmpB, PmpC, PmpF and PmpH, as described further below:

PmpA Fusion Proteins

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PmpA is 107 kD protein containing 982 aa and was cloned from serovar E. The PmpA protein was divided into 2 overlapping fragments, the PmpA(N-terminal) and (C-terminal) portions.

PmpA(N-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCATGTTTATAACAAAGGAACTTATG (SEQ ID NO: 306)

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GAGAGCGGCCGCTTACTTAGGTGAGAAGAAGGGAGTTTC (SEQ ID NO: 307) respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 308, encoding a 66 kD protein (619aa) expressing the segment 1-473 aa of PmpA. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 309.

PmpA(C-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCCATTCTATTCATTTCTTTGATCCTG (SEQ ID NO: 310)

GAGAGCGGCCGCTTAGAAGCCAACATAGCCTCC (SEQ ID NO: 311)

respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 312, encoding a 74 kD protein (691aa) expressing the segment 438-982 aa of PmpA. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 313.

PmpF Fusion Proteins

PmpF is 112 kD protein containing 1034 aa and was cloned from the serovar E. PmpF protein was divided into 2 overlapping fragments, the PmpF(N- term) and (C-term) portions.

PmpF(N-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCATGATTAAAAGAACTTCTCTATCC (SEQ ID NO: 314)

GAGAGCGGCCGCTTATAATTCTGCATCATCTTCTATGGC (SEQ ID NO: 315)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 316, encoding a 69 kD protein (646aa) expressing the segment 1-499 aa of PmpF. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 317.

PmpF(C-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGACATACGAACTCTGATGGG (SEQ ID NO: 318)

GAGAGCGGCCGCTTAAAAGACCAGAGCTCCTCC (SEQ ID NO: 319)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 320, encoding a 77 kD protein (715aa) expressing the segment 466-1034aa of PmpF. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 321.

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PmpH Fusion Proteins

PmpH is 108 kD protein containing 1016 as and was cloned from the serovar E. PmpH protein was divided into 2 overlapping fragments, the PmpH(N-term)and (C-term)portions.

PmpH(N-term) was amplified by the sense and antisense primers:

GAGAGCGCCGCTCATGCCTTTTTCTTTGAGATCTAC (SEQ ID NO: 322) GAGAGCGCCGCTTACACAGATCCATTACCGGACTG (SEQ ID NO: 323)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 324, encoding a 64 kD protein (631aa) expressing the segment 1-484 aa of PmpH. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 325. The donor line CHH037 was found to be reactive against this protein.

PmpH(C-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGATCCTGTAGTACAAAATAATTCAGC (SEQ ID NO: 326) GAGAGCGGCCGCTTAAAAGATTCTATTCAAGCC (SEQ ID NO: 327)

15 respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 328, encoding a 77 kD protein (715aa) expressing the segment 449-1016aa of PmpH. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 329. The patient line CT12 was found to be reactive in response to this protein.

PmpB Fusion Proteins

PmpB is 183 kD protein containing 1750 aa and was cloned from the serovar E. PmpB protein was divided into 4 overlapping fragments, PmpB(1), (2), (3) and (4).

PmpB(1) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCATGAAATGGCTGTCAGCTACTGCG (SEQ ID NO: 330)

25 GAGAGCGGCCGCTTACTTAATGCGAATTTCTTCAAG (SEQ ID NO: 331)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 332, and encodes is a 53 kD protein (518aa) expressing the segment 1-372 aa of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 333.

PmpB(2) was amplified by the sense and antisense primers:

GAGAGCGCCCCCCGGTGACCTCTCAATTCAATCTTC (SEQ ID NO: 334)

GAGAGCGGCCGCTTAGTTCTCTGTTACAGATAAGGAGAC (SEQ ID NO: 335)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 336 and encodes a 60 kD protein (585aa) expressing the segment 330-767 aa of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 337. Cell lines derived from patient lines CT1, CT3, CT4 responded to this recombinant pmpB protein.

PmpB(3) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGACCAACTGAATATCTCTGAGAAC (SEQ ID NO: 338)

10 GAGCGGCCGCTTAAGAGACTACGTGGAGTTCTG (SEQ ID NO: 339)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 340 encodes a 67 kD protein (654aa) expressing the segment 732-1236 aa of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 341

PmpB(4) was amplified by the sense and antisense primers:

15 GAGAGCGCCCCTCGGAACTATTGTGTTCTCTTCTG (SEQ ID NO: 342)

GAGAGCGCCGCTTAGAAGATCATGCGAGCACCGC (SEQ ID NO: 343)

respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 344 encodes a 76 kD protein (700aa) expressing the segment 1160-1750 of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 345.

20 PmpC Fusion Proteins

PmpC is 187 kD protein containing 1774 aa and was cloned from the serovar E/L2. PmpC protein was divided into 3 overlapping fragments, PmpC(1), (2) and (3).

PmpC(1) was amplified by the sense and antisense primers:

25 GAGAGCGCCGCTCATGAAATTTATGTCAGCTACTGC (SEQ ID NO: 346)
GAGAGCGGCCGCTTACCCTGTAATTCCAGTGATGGTC (SEQ ID NO: 347)

respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 348 and encodes a 51 kD protein (487aa) expressing the segment 1-340 aa of PmpC. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 349.

PmpC(2) was amplified by the sense and antisense primers:

- 5 GAGAGCGGCCGCTCGATACACAAGTATCAGAATCACC (SEQ ID NO: 350)
 GAGAGCGGCCGCTTAAGAGGACGATGAGACACTCTCG (SEQ ID NO: 351)
 respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 352 and encodes a 60 kD protein (583aa) expressing the segment 305-741 aa of PmpC. The amino acid sequence of the fusion protein is set forth in SEO ID NO: 353.
 - PmpC(3) was amplified by the sense and antisense primers:

 GAGAGCGGCCGCTCGATCAATCTAACGAAAACACAGACG (SEQ ID NO: 354)

 GAGAGCGGCCGCTTAGACCAAAGCTCCATCAGCAAC (SEQ ID NO: 355)

 respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 356 and encodes a 70 kD protein (683aa) expressing the segment 714-1250 aa of PmpC. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 357.

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EXAMPE 11

IMMUNOGENICITY OF CT622

Chlamydia-specific T cells lines were generated from two patients with Chlamydia infections and the lines were designated CT1 and CT13. The T cell lines were either generated against monocyte-derived dendritic cells infected *C. trachomatis* serovar E for 72 hours (CT1-ERB) or against killed serovar E elementary bodies (EB) (CT13-EEB). Once generated, the lines were tested against the recombinant Chlamydia-specific protein, CT622 in a proliferation assay. Proliferation assays were performed by stimulating 2.5x10⁴ T cells in the presence of 1x10⁴ monocyte-derived dendritic cells with either recombinant CT antigens (2µg/ml) or Chlamydia EBs (1µg/ml). The assay was incubated for 4 days with a ³H-thymidine pulse for the last 18 hours.

134

The cell line CT1-ERB demonstrated proliferative responses significantly above the media controls when stimulated with CT622, CT875, and CT EB. The cell line CT13-EEB demonstrated a proliferative response significantly above media controls when stimulated with CT622, CT875, and CT EB (see Figure 12).

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EXAMPLE 12

CLONING AND EXPRESSION OF FULL LENGTH CHLAMYDIA TRACHOMATIS GENES CT611, ORF3 AND OppA1

Recombinant protein expression of the full-length open reading frames was performed for clones containing genes CT611, ORF-3, and oppA1. The clones that contained the genes of interest were CtL2-8 (SEQ ID NO:285) which encoded 4 ORFs (CT474, CT473, CT060, and CT139), CtL2-10 (SEQ ID NO:284) which encoded the ORFs of CT610 and CT611, and clones 16CtL2-16 (SEQ ID NO:47), 16-D4-22 (SEQ ID NO:119) and 19-A5-54 (SEQ ID NO:84) which all contained sequences related to ORF-3. Sequences within CtL2-10 (Ct-610) and CtL2-16 (ORF-3) were also independently identified by the T-cell expression cloning approach. The clone CtL2-8 was further investigated as this clone had stimulated the proliferative responses and IFN-gamma production by two T cell lines generated against serovar E.

Cloning and expression of clone sequences:

CtL2-10 was found to encode two open reading frames (ORFs), CT610 and CT611, and these were found organized adjacent to each other within the genomic clone. The full length ORF of CT610 (containing a PQQ synthesis domain) was previously expressed and demonstrated to stimulate the proliferative responses of T cell lines generated against Chlamydia. To determine whether the second ORF, CT611, was also recognized by T cells, the full-length sequence of CT611 was PCR amplified and engineered for protein expression. The nucleotide sequence is disclosed in SEQ ID NO:361 with the corresponding amino acid sequence disclosed in SEQ ID NO:365.

The second serological clone, CtL2-8, was found to contain 4 ORFs (CT474, CT473, CT060, and CT139). Overlapping peptides to the three smallest predicted ORFs (CT474, CT473, and CT060) did not stimulate the proliferative responses of T

cell lines. This suggested that the immunostimulatory antigen resides in the fourth ORF, CT139. The ORF of CT139 is approximately 450 nucleotides. The full-length nucleotide sequence is disclosed in SEQ ID NO:359 and the full-length amino acid sequence is disclosed in SEQ ID NO:363. Amino acid sequence comparison from Genbank revealed that it is an oligo-peptide binding protein (oppA1) as well as belonging to the peptide ABC transporter family. This protein is 462 amino acids long with a predicted size of 48.3kDa and appears to contain 2 trans-membrane regions.

To express the full-length sequence of oppA1, oligonucleotides were designed which specifically amplified sequences starting from amino acid residue 22 (devoid of the first transmembrane domain), the nucleotide sequence for which is disclosed in SEQ ID NO:358 and, the amino acid sequence of which is disclosed in SEQ ID NO:362. This was shown to express the protein in E. coli.

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The full-length cloning and recombinant protein expression of ORF-3 was also achieved. The nucleotide and amino acid sequences are disclosed in SEQ ID NOs:360 and 364, respectively.

EXAMPLE 13 RECOMBINANT CHLAMYDIAL ANTIGENS RECOGNIZED BY T CELL LINES

Patient T cell lines were generated from the following donors: CT1, CT2, CT3, CT4, CT5, CT6, CT7, CT8, CT9, CT10, CT11, CT12, CT13, CT14, CT15, and CT16, some of which were discussed above. A summary of their details is included in Table V.

	Table V: C. trachomatis patients											
Patients	Gender Age		Clinical Manifestation	Serovar	IgG titer	Multiple Infections						
CT1	M	27	NGU	LCR	Negative	No						
CT2	M	24	NGU	D	Negative	E						
CT3	М	43	Asymptomatic	J	Ct 1:512	No						

PCT/US01/23121

Shed Eb					Υ		т
CT4 F 25 Asymptomatic Shed Eb J Ct Y CT5 F 27 BV LCR Ct 1:256 F/F CT6 M 26 Perinial rash Discharge, dysuria G Cp 1:1024 N CT7 F 29 BV Genital ulcer E Ct 1:512 N Cp 1:1024 N CT8 F 24 Not Known LCR Not kested NA tested CT9 M 24 asymptomatic LCR Ct 1:128 N Cp 1:128 N CT10 F 20 Mild itch vulvar negative negative negative negative negative plib 6/96 12/1/98 CT11 F 21 BV Abnormal pap smear J Ct 1:512 F/F/J/E/E PID 6/96 CT12 M 20 asymptomatic LCR Cp 1:512 N N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct vaginal discharge, dysuria				Shed Eb		Ср	
CT4 F 25 Asymptomatic Shed Eb J Ct Y CT5 F 27 BV LCR Ct 1:256 F/F CT6 M 26 Perinial rash Discharge, dysuria G Cp N 1:1024 CT7 F 29 BV E Ct 1:512 N CT8 F 24 Not Known LCR Not NA tested CT9 M 24 asymptomatic LCR Ct 1:128 N CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear Smear Ct ct Ct N 1:512 CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N			-	Dx was HPV		1:1024	
CT4 F 25 Asymptomatic Shed Eb J Ct Y CT5 F 27 BV LCR Ct 1:256 F/F CT6 M 26 Perinial rash Discharge, dysuria G Cp 1:1024 N CT7 F 29 BV E Ct 1:512 N CT8 F 24 Not Known LCR Not cycle NA CT9 M 24 asymptomatic LCR Ct 1:128 N CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear Smear Ct 20 N 1:512 N CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N						Cps	
CT5						1:256	
CT5 F 27 BV LCR Ct 1:256 F/F CT6 M 26 Perinial rash Discharge, dysuria G Cp 1:1024 N CT7 F 29 BV Genital ulcer E Ct 1:512 N N CT8 F 24 Not Known LCR Not tested NA tested CT9 M 24 asymptomatic LCR Ct 1:128 N N CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV Johnormal pap smear J Ct 1:512 F/F/I/E/E PID 6/96 CT12 M 20 asymptomatic LCR Cp N N 1:512 CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N	CT4	F	25	Asymptomatic	J	Ct	Y
CT6 M 26 Perinial rash Discharge, dysuria G Cp 1:256 N CT7 F 29 BV Genital ulcer E Ct 1:512 N N CT8 F 24 Not Known LCR Not tested NA NA CT9 M 24 asymptomatic LCR Ct 1:128 N N CT9 M 24 asymptomatic LCR Ct 1:128 N N CT9 M 20 Mild itch vulvar negative negative negative negative PiD 6/96 12/1/98 CT10 F 20 Mild itch vulvar negative negative negative PiD 6/96 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E PID 6/96 CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N				Shed Eb		1:1024	
CT6 M 26 Perinial rash Discharge, dysuria G Cp 1:256 N CT7 F 29 BV Genital ulcer E Ct 1:512 N N CT8 F 24 Not Known LCR Not tested NA NA CT9 M 24 asymptomatic LCR Ct 1:128 N N CT9 M 24 asymptomatic LCR Ct 1:128 N N CT9 M 20 Mild itch vulvar negative negative negative negative negative negative Abnormal pap smear D Ct 1:512 F/F/J/E/E PID 6/96 CT12 M 20 asymptomatic LCR Cp N N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N							
CT6 M 26 Perinial rash Discharge, dysuria G Cp N 1:1024 N CT7 F 29 BV E Ct 1:512 N Cp 1:1024 Not Cp 1:1024 Not Cp 1:1024 CT8 F 24 Not Known LCR Not tested NA tested NA Cp 1:128 N Cp 1:128 CT9 M 24 asymptomatic LCR Ct 1:128 N Cp 1:128 N Cp 1:128 CT10 F 20 Mild itch vulvar negative negative negative negative plD 6/96 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E PID 6/96 CT12 M 20 asymptomatic LCR Cp N 1:512 N 1:512 CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N 1:1024	CT5	F	27	BV	LCR	Ct 1:256	F/F
CT6 M 26 Perinial rash Discharge, dysuria G Cp 1:1024 N CT7 F 29 BV E Ct 1:512 N CT8 F 24 Not Known LCR Not Known tested NA CT9 M 24 asymptomatic LCR Ct 1:128 N CT9 I 20 Mild itch vulvar negative 12/1/98 CT10 F 20 Mild itch vulvar negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear LCR Cp N CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N	}	Ì		1		Ср	
Discharge, dysuria						1:256	
CT7	CT6	M	26	Perinial rash	G	Ср	N
CT7 F 29 BV E Ct 1:512 N Cp 1:1024 Cp 1:1024 NA CT8 F 24 Not Known LCR Not kested NA CT9 M 24 asymptomatic LCR Ct 1:128 N Cp 1:128 Cp 1:128 CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear Smear LCR Cp N CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N				Discharge,		1:1024	
CT8 F 24 Not Known LCR Not NA tested				dysuria			
CT8 F 24 Not Known LCR Not tested CT9 M 24 asymptomatic LCR Ct 1:128 N CT9 M 24 asymptomatic LCR Ct 1:128 N CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear Abnormal pap smear LCR Cp N CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N	CT7	F	29	BV	Е	Ct 1:512	N
CT8 F 24 Not Known LCR Not tested CT9 M 24 asymptomatic LCR Ct 1:128 N Cp 1:128 CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV Abnormal pap smear CT12 M 20 asymptomatic LCR Cp 1:512 F/F/J/E/E PID 6/96 CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria				Genital ulcer		Cp	
CT9 M 24 asymptomatic LCR Ct 1:128 N CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear PID 6/96 CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N		İ				1:1024	
CT9 M 24 asymptomatic LCR Ct 1:128 N Cp 1:128 CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear CT12 M 20 asymptomatic LCR Cp 1:512 CT13 F 18 BV, gonorrhea, G Ct Ct vaginal discharge, dysuria	CT8	F	24	Not Known	LCR	Not	NA
CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear CT12 M 20 asymptomatic LCR Cp N 1:512 CT13 F 18 BV, gonorrhea, G Ct N Ct vaginal discharge, dysuria						tested	
CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear Abnormal pap smear PID 6/96 CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N	СТ9	M	24	asymptomatic	LCR	Ct 1:128	N
CT10 F 20 Mild itch vulvar negative negative 12/1/98 CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear CT12 M 20 asymptomatic LCR Cp N 1:512 CT13 F 18 BV, gonorrhea, G Ct N Ct vaginal discharge, dysuria						Cp	
CT11 F 21 BV J Ct 1:512 F/F/J/E/E Abnormal pap smear LCR Cp N N CT12 M 20 asymptomatic LCR Cp N CT13 F 18 BV, gonorrhea, Ct vaginal discharge, dysuria G Ct N						1:128	
Abnormal pap smear CT12 M 20 asymptomatic LCR Cp N 1:512 CT13 F 18 BV, gonorrhea, G Ct N 1:1024 discharge, dysuria	CT10	F	20	Mild itch vulvar	negative	negative	12/1/98
CT12 M 20 asymptomatic LCR Cp N 1:512 CT13 F 18 BV, gonorrhea, G Ct N 1:1024 discharge, dysuria	CT11	F	21	BV	J	Ct 1:512	F/F/J/E/E
CT12 M 20 asymptomatic LCR Cp N 1:512 CT13 F 18 BV, gonorrhea, G Ct N Ct vaginal discharge, dysuria				Abnormal pap			PID 6/96
CT13 F 18 BV, gonorrhea, G Ct N Ct vaginal discharge, dysuria				smear			
CT13 F 18 BV, gonorrhea, G Ct N Ct vaginal discharge, dysuria	CT12	M	20	asymptomatic	LCR	Ср	N
Ct vaginal 1:1024 discharge, dysuria						1:512	
discharge, dysuria	CT13	F	18	BV, gonorrhea,	G	Ct	N
				Ct vaginal		1:1024	
CT14 M 24 NGU LCR Ct 1:256 N				discharge, dysuria			
	CT14	M	24	NGU	LCR	Ct 1:256	N

					Cp 1:256	
CT15	F	21	Muco-purulint cervicitis Vaginal discharge	culture	Ct 1:256 Ct IgM 1:320 Cp 1:64	N
CT16	М	26	Asymptomatic/ contact	LCR	NA	N
CL8	М	38	No clinical history of disease	negative	negative	N

NGU=Non-Gonococcal Urethritis; BV=Bacterial Vaginosis; CT=Chlamydia trachomatis; Cp=Chlamydia pneumoniae; Eb=Chlamydia elementary bodies; HPV=human papiloma virus; Dx=diagnosis; PID=pelvic inflammatory disease; LCR=Ligase chain reaction.

PBMC were collected from a second series of donors and T cell lines have been generated from a sub-set of these. A summary of the details for three such T cell lines is listed in the table below.

10

15

	Table III: Normal Donors											
Donor	Gender	Age	CT IgG Titer	CP IgG Titer								
CHH011	F	49	1:64	1:16								
СНН037	F	22	0	0								
CHH042	F	25	0	1:16								

Donor CHH011 is a heathly 49 year old female donor sero-negative for C. trachomatis. PBMC produced higher quantities of IFN-gamma in response to C. trachomatis elementary bodies as compared to C. pneumoniae elementary bodies, indicating a C. trachomatis-specific response. Donor CHH037 is a 22 year old healthy

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WO 02/08267

10

15

138

PCT/US01/23121

female donor sero-negative for C. trachomatis. PBMC poruced higher quantities of IFN-gamma in response to C. trachomatis elementary bodies as compared to C. pneumoniae elementary bodies, indicating a C. trachomatis-specific response. CHH042 is a 25 year old healthy female donor with an IgG titer of 1:16 to C. pneumoniae. PBMC produced higher quantities of IFN-gamma in response to C. trachomatis elementary bodies as compared to C. pneumoniae elementary bodies, indicating a C. trachomatis-specific response.

Recombinant proteins for several *Chlamydia trachomatis* genes were generated as described above. Sequences for MOMP was derived from serovar F. The genes CT875, CT622, pmp-B-2, pmpA, and CT529 were derived from serovar E and sequences for the genes gro-EL, Swib, pmpD, pmpG, TSA, CT610, pmpC, pmpE, S13, lpdA, pmpI, and pmpH-C were derived from LII.

Several of the patient and donor lines described above were tested against the recombinant Chlamydia proteins. Table IV summarizes the results of the T cell responses to these recombinant Chlamydia proteins.

	Table	VII: R	ecoml	binant	Chlam	ydia An	tigens I	Recogni	zed By	T Cel	1 Lines	3	
Antigen	Sero-	#of	С	СТ	CT1	CT3	CT4	CT5	СТ	CT	СТ	СН	CH
	var	hits	L8	10	E	E	L2	E	11	12	13	H-	H-
			L2	E					Е	E	E	011	037
						<u> </u>						Е	Е
gro-EL (CT110)	L2	10	1	+	+	+	+	+	+	+	+	+	+
MompF (CT681)	F	10	-	+	+	+	+	+	+	+	+	+	+
CT875	E	8	-	+	+	-	+	+	+	+	+	-	+
SWIB (CT460)	L2	8	+	+	<u>-</u>	+	-	+	1	+	+	+	+
pmpD (CT812)	L2	5	-	+	+	+	+	-	1	+	+	1	-

139

pmpG	L2	6	-	+	+	_	+	+	nt	-	+	+	_
(CT871)													
TSA	L2	6	-	-	+	+	+	+	-	-	+	-	+
(CT603)													
CT622	E	3	-	-	+	-	+	-	-	-	+	-	-
CT610	L2	3	-	+	-	+	•	-	-	+	-	~	-
pmpB-2	Е	3	-	-	+	+	+	_	-	-	-	1	-
(CT413)													:
pmpC	L2	4	-	-	-	+	-	+	-	+	-	1	+
(CT414)						j						_	
pmpE	L2	3	•	+	+	-	•	-	-	•	+	-	-
(CT869)													
S13	L2	2	+	-	-	-	+	-	-	-	-	-	-
(CT509)				:									
lpdA	L2	3	-	-	+	+	-	-	-	-	-	+	-
(CT557)													
pmpI	L2	2	-	-	+	-	-	-	-	-	-	+	-
(CT874)													
pmpH-C	L2	1	-	-	-	-	-	-	-	+	-	-	-
(CT872)													
pmpA	E	0	-	-	-	-	-	-	-	-	-	-	-
(CT412)													
CT529	Е	0	-	-	-	-	-	-	-	-	•	-	-
		l				L							

Although the present invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, changes and modifications can be carried out without departing from the scope of the invention which is intended to be limited only by the scope of the appended claims.

140

Claims

What is Claimed:

- 1. A composition for eliciting an immune response comprising a Chlamydia Cap1 protein or an immunogenic fragment thereof and an immunostimulant.
- 2. The composition of claim 1, wherein the immunogenic fragment comprises at least a CTL epitope consisting essentially of amino acids 139-147 of a Cap1 protein.
- 3. The composition of claim 1, wherein the Cap 1 protein comprises an amino acid sequence set forth in SEQ ID NO: 121 or a sequence having at least about 90% identity to the sequence set forth in SEQ ID NO: 121.
- 4. The composition of claim 1, wherein the Cap1 protein or immunogenic fragment thereof comprises a sequence set forth in any one of SEQ ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.
- 5. The composition of claim 1, wherein the immunogenic fragment comprises amino acids 107-176 of a Cap1 protein.
- 6. The composition of claim 5, wherein the immunogenic fragment comprises amino acids 107-176 of a Cap1 protein having an amino acid sequence set forth in any one of SEQ ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.
 - 7. The composition of claim 1, wherein the immunogenic fragment is

WO 02/08267

immunologically reactive with a CD8+ T-cell of a Chlamydia-infected animal.

- 8. A method for stimulating a Chlamydia-specific T-cell response in an animal comprising administering to an animal an effective amount of a composition according to claim 1.
- 9. A method for inhibiting the development of a Chlamydia infection in an animal, comprising administering to an animal an effective amount of a composition according to claim 1.
- 10. A composition for eliciting an immune response comprising an isolated polynucleotide that encodes a Chlamydia Cap1 protein or an immunogenic fragment thereof and an immunostimulant.
- 11. The composition of claim 10, wherein the immunogenic fragment comprises at least the CTL epitope sequence consisting essentially of amino acids 139-147 of a Cap1 protein.
- 12. The composition of claim 10, wherein the Cap 1 protein comprises an amino acid sequence set forth in SEQ ID NO: 121 or a sequence having at least about 90% identity to the sequence set forth in SEQ ID NO: 121.
- 13. The composition of claim 10, wherein the Cap1 protein or immunogenic fragment thereof comprises a sequence set forth in any one of SEQ ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.

- 14. The composition of claim 10, wherein the immunogenic fragment comprises amino acids 107-176 of a Cap1 protein.
- 15. The composition of claim 14, wherein the immunogenic fragment comprises amino acids 107-176 of a Cap1 protein having an amino acid sequence set forth in any one of SEQ ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.
- 16. The composition of claim 10, wherein the immunogenic fragment is immunologically reactive with a CD8+ T-cell of a Chlamydia-infected animal.
- 17. The composition of claim 10, wherein the isolated polynucleotide is operably linked within a viral delivery vector.
- 18. The composition of claim 17, wherein the viral delivery vector is a vaccinia virus delivery vector.
- 19. A method for stimulating a Chlamydia-specific T-cell response in an animal comprising administering to said animal an effective amount of a composition according to claim 10.
- 20. A method for inhibiting the development of a Chlamydia infection in an animal, comprising administering to an animal said effective amount of a composition according to claim 10.

- 21. A method for inhibiting the development of a Chlamydia infection in an animal, comprising administering to said animal an effective amount of a composition according to claim 18.
- 22. An isolated polynucleotide comprising a sequence selected from the group consisting of:
 - (a) sequences provided in SEQ ID NO:358-361;
 - (b) complements of the sequences provided in SEQ ID NO:358-361;
- (c) sequences consisting of at least 20 contiguous residues of a sequence provided in SEQ ID NO:358-361;
- (d) sequences that hybridize to a sequence provided in SEQ ID NO:358-361, under highly stringent conditions;
- (e) sequences having at least 95% identity to a sequence of SEQ ID NO:358-361;
- (f) sequences having at least 99% identity to a sequence of SEQ ID NO:358-361; and
- (g) degenerate variants of a sequence provided in SEQ ID NO:358-361.
- 23. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of:
 - (a) sequences encoded by a polynucleotide of claim 22;
- (b) sequences having at least 95% identity to a sequence encoded by a polynucleotide of claim 22; and
- (c) sequences having at least 99% identity to a sequence encoded by a polynucleotide of claim 22.
- 24. An isolated polypeptide comprising at least an immunogenic fragment of a polypeptide sequence selected from the group consisting of:
 - (a) a polypeptide sequence set forth in SEQ ID NO:362-365,
- (b) a polypeptide sequence having at least 95% identity with a sequence set forth in SEQ ID NO:362-365, and

(c) a polypeptide sequence having at least 99% identity with a sequence set forth in SEQ ID NO:362-365.

- 25. An expression vector comprising a polynucleotide of claim 22 operably linked to an expression control sequence.
- 26. A host cell transformed or transfected with an expression vector according to claim 25.
- 27. An isolated antibody, or antigen-binding fragment thereof, that specifically binds to a polypeptide of any one of claims 23 and 24.
- 28. A method for detecting the presence of Chlamydia in a patient, comprising the steps of:
 - (a) obtaining a biological sample from the patient;
- (b) contacting the biological sample with a binding agent that binds to a polypeptide of any one of claims 23 and 24;
- (c) detecting in the sample an amount of polypeptide that binds to the binding agent; and
- (d) comparing the amount of polypeptide to a predetermined cut-off value and therefrom determining the presence of Chlamydia in the patient.
- 29. A fusion protein comprising at least one polypeptide according to claim 23 or claim 24.
- 30. An oligonucleotide that hybridizes to a sequence recited in SEQ ID NO: 358-361 under highly stringent conditions.
- 31. A method for stimulating and/or expanding T cells specific for a Chlamydia protein, comprising contacting T cells with at least one component selected from the group consisting of:
 - (a) a polypeptide according to claim 23 or claim 24;
 - (b) a polynucleotide according to claim 22; and

(c) an antigen-presenting cell that expresses a polynucleotide according to claim 22,

under conditions and for a time sufficient to permit the stimulation and/or expansion of T cells.

- 32. An isolated T cell population, comprising T cells prepared according to the method of claim 31.
- 33. A composition comprising a first component selected from the group consisting of physiologically acceptable carriers and immunostimulants, and a second component selected from the group consisting of:
 - (a) a polypeptide according to claim 23 or claim 24;
 - (b) a polynucleotide according to claim 22;
 - (c) an antibody according to claim 27;
 - (d) a fusion protein according to claim 29;
 - (e) a T cell population according to claim 32; and
- (f) an antigen presenting cell that expresses a polypeptide according to claim 23 or claim 24.
- 34. A method for stimulating an immune response in a patient, comprising administering to the patient a composition selected from the group consisting of;
 - (a) a composition of claim 33;
 - (b) a polynucleotide sequence of any one of SEQ ID NO:407-430, 525-559, and 582-598; and
 - (c) a polypeptide sequence of any one of SEQ ID NO:431-454 and 560-581.
- 35. A method for the treatment of Chlamydia infection in a patient, comprising administering to the patient a composition selected from the group consisting of;
 - (a) a composition of claim 33;

- (b) a polynucleotide sequence of any one of SEQ ID NO: 407-430, 525-559, and 582-598; and
- (d) a polypeptide sequence of any one of SEQ ID NO: 431-454 and 560-581.
- 36. A method for determining the presence of Chlamydia in a patient, comprising the steps of:
 - (a) obtaining a biological sample from the patient;
- (b) contacting the biological sample with an oligonucleotide according to claim 30;
- (c) detecting in the sample an amount of a polynucleotide that hybridizes to the oligonucleotide; and
- (d) comparing the amount of polynucleotide that hybridizes to the oligonucleotide to a predetermined cut-off value, and therefore determining the presence of the cancer in the patient.
- 37. A diagnostic kit comprising at least one oligonucleotide according to claim 30.
- 38. A diagnostic kit comprising at least one antibody according to claim 27 and a detection reagent, wherein the detection reagent comprises a reporter group.
- 39. A method for the treatment of Chlamydia in a patient, comprising the steps of:
- (a) incubating CD4+ and/or CD8+ T cells isolated from a patient with at least one component selected from the group consisting of:
 - (i) a polypeptide according to any one of claims 23 and 24;
 - (ii) a polypeptide sequence of any one of SEQ ID NO: 431-454 and 560-581;
 - (iii) a polynucleotide according to claim 22;
 - (iv) a polynucleotide sequence of any one of SEQ ID NO: 407-430, 525-559 and 582-598;

- (v) an antigen presenting cell that expresses a polypeptide sequence set forth in any one of claims 23 and 24;
- (vi) an antigen presenting cell that expresses a polypeptide sequence of any one of SEQ ID NO: 431-454 and 560-581, such that the T cells proliferate; and
- (b) administering to the patient an effective amount of the proliferated T cells.

1/11

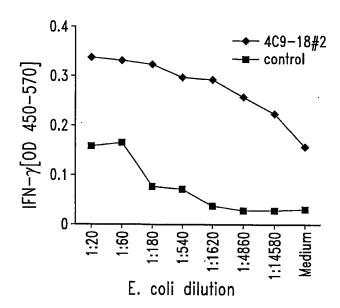


Fig. 1

2/11

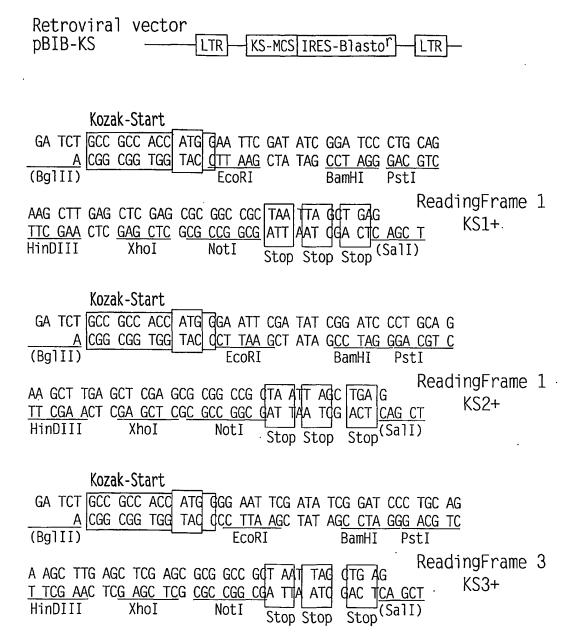


Fig. 2

Chlamydia C17.8 Peptide Screen

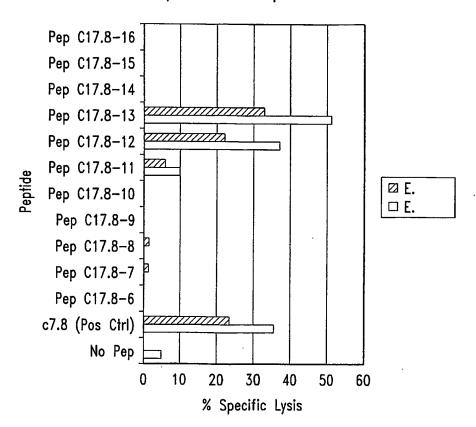
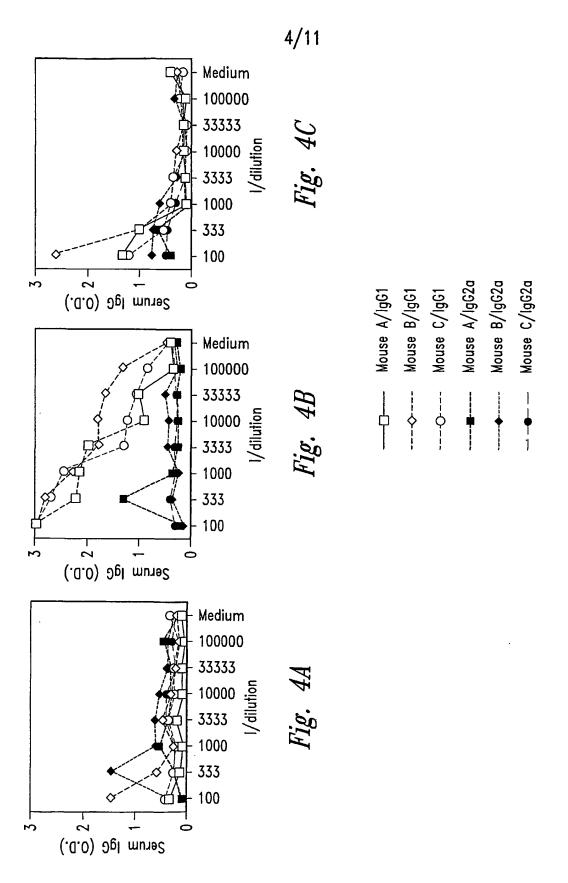
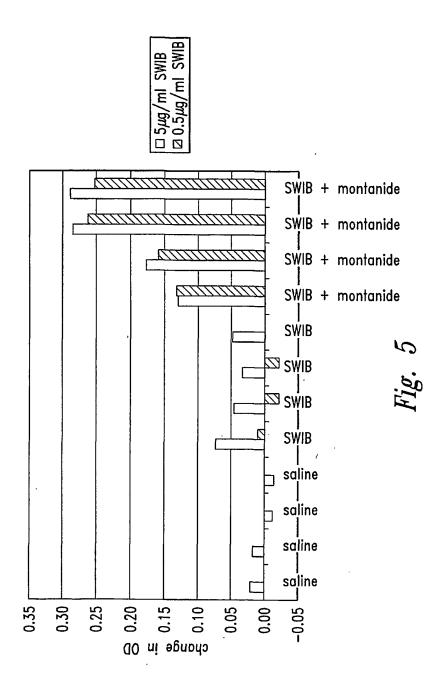


Fig. 3

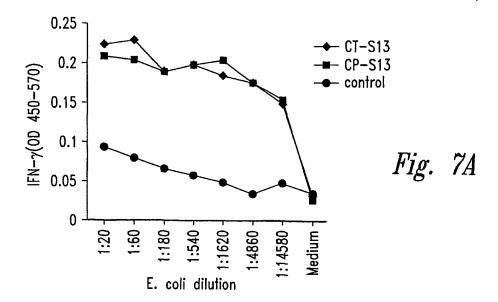


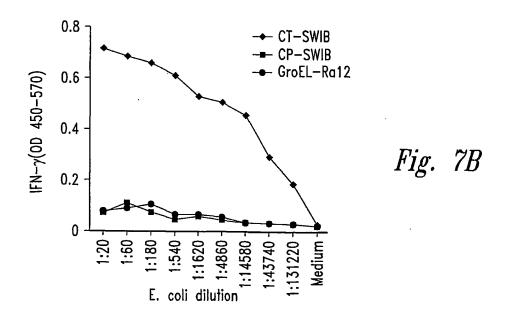


6/11

- CP SWIB Nde (5' primer)
- CP SWIB EcoRI (3' primer)
- 5' CTCGAGGAATTCTTATTTTACAATATGTTTGGA
- CP S13 Nde (5' primer)
- 5' GATATACATATGCATCACCATCACCATCACATGCCACGCATCATTGGAATGAT
- CP S13 EcoRI (3' primer)
 5' CTCGAGGAATTCTTATTTCTTCTTACCTGC

Fig. 6





8/11

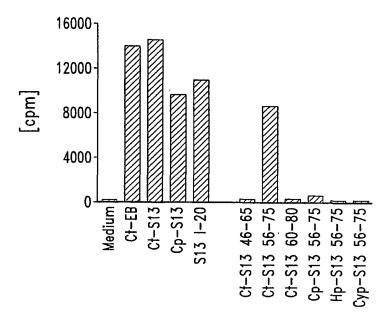
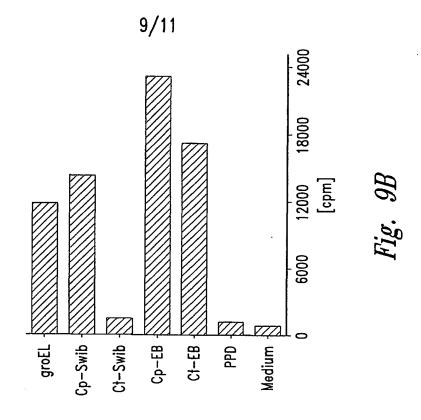
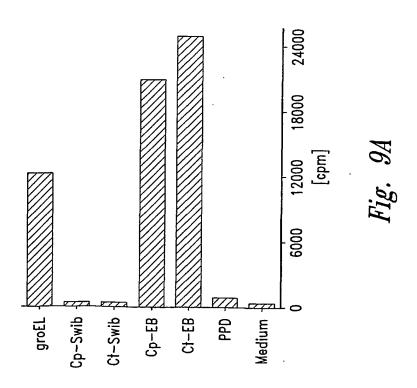
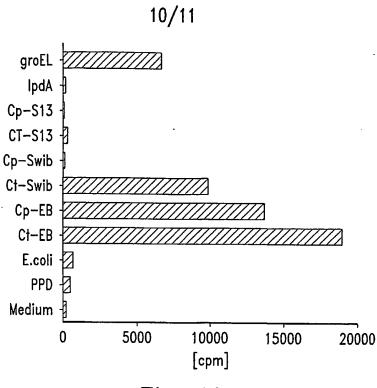


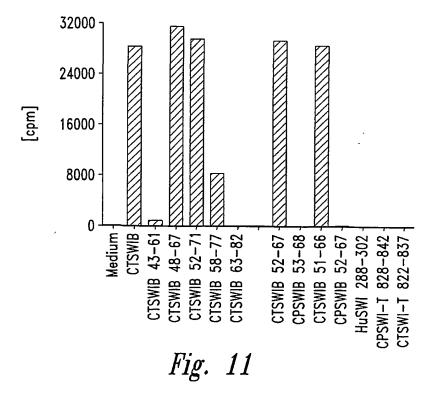
Fig. 8

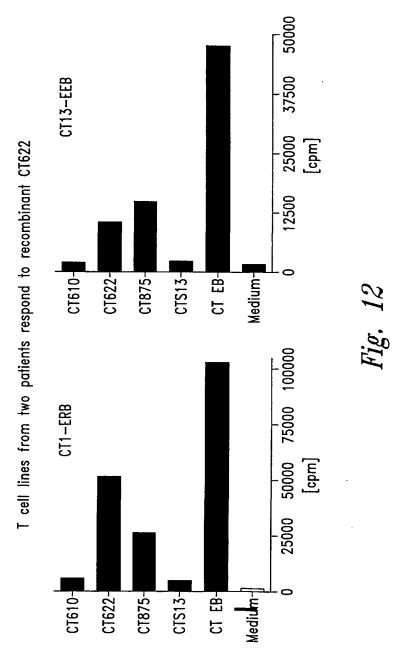












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 Arg Phe Phe Leu Pro Lys Leu Lys Gln Ile Trp Asp Leu Leu Ala
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 Ile Leu Trp Arg Leu Thr Met Gln Arg Leu Trp Trp Val Leu Asp Ser
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 Leu Ser Val Arg Lys Glu Gln Ile Ala Lys Pro Ala Ala Leu Val Leu
 Arg Glu Lys Ser Arg Tyr Ser Lys Cys Arg Glu Arg Lys Met Leu Ala
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                                105
                                                     110
Arg Arg Lys Ser Leu Glu Arg Lys Pro Arg Arg Ser Arg Ala Ser Ser
115 120 125
                                               125
 Met His Ser Ser Leu Cys Ser Arg Ser Phe Trp Asn Ala Leu Pro Thr
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Phe Ser Asn Trp Cys Arg Cys Leu Leu Gln Trp Val Phe Val Arg Leu 145 150 150 160
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 Trp Leu Leu Asp Val Arg Ser Leu Leu Gln Leu Leu Asp Cys Ala Leu
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                                    170
 Ser Ala Pro Glu His Lys Gly Phe Phe Lys Phe Leu Lys Lys Lys Ala
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 Val Ser Lys Lys Lys Gln Pro Phe Leu Ser Thr Lys Cys Leu Ala Phe
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                                                                        180
 teaccegaca aggacteegt ttegtactag aageetetgt ateaaatatt gaggatatag
                                                                        240
gagatogogt toggttaact atcaatggga atgtogaaga atacgattac gttotogtat
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                                                                        420
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tctataggac gccqtttgaa taCagaaaat attggcttgg ataaagctgg tgttatttgt
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                                                                      540
gtgatettta cettecetga agtegettea gtaggeetet eeccaacage ageteaacaa
                                                                      600
catctccttc ttcgcttact ttttctgaaa aatttgatac agaagaagaa ttcctcgcac
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                            40
                                              45
Asn Asn Pro Asp Ile Ser Lys Thr Met Phe Asp Lys Phe Thr Arg Gln
                        55
                                            60
Gly Leu Arg Phe Val Leu Glu Ala Ser Val Ser Asn Ile Glu Asp Ile
                    70
                                        75
Gly Asp Arg Val Arg Leu Thr Ile Asn Gly Asn Val Glu Glu Tyr Asp
                                    90
Tyr Val Leu Val Ser Ile Gly Arg Arg Leu Asn Thr Glu Asn Ile Gly
           100
                               105
Leu Asp Lys Ala Gly Val Ile Cys Asp Glu Arg Gly Val Ile Pro Thr
       115
                            120
                                                125
Asp Ala Thr Met Arg Thr Asn Val Pro Asn Ile Tyr Ala Ile Gly Asp
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                                            140
Ile Thr Gly Lys Trp Gln Leu Ala His Val Ala Ser His Gln Gly Ile
Ile Ala Ala Arg Asn Ile Gly Gly His Lys Glu Glu Ile Asp Tyr Ser
                                    170
                                                        175
               165
Ala Val Pro Ser Val Ile Phe Thr Phe Pro Glu Val Ala Ser Val Gly
                                                    190
                                185
           180
Leu Ser Pro Thr Ala Ala Gln Gln His Leu Leu Leu Arg Leu Leu Phe
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                           200
                                                205
Leu Lys Asn Leu Ile Gln Lys Lys Asn Ser Ser His Thr Cys Glu Glu
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tacattaaaa aacacaactg tcaggatcaa aaaaataaac gtaatatcct tcccgatgcg
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<211> 87

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                            40
                                                45
Asp Gln Lys Asn Lys Arg Asn Ile Leu Pro Asp Ala Asn Leu Ala Lys
                        55
Val Phe Gly Ser Ser Asp Pro Ile Asp Met Phe Gln Met Thr Lys Ala
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cctgaggcaa gagcctctga attaactgaa gaagaagtag gacgactgaa ctctctgcta
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caatcagaat ataccgtaga aggggatttg cgacgtcgtg ttcaatcgga tatcaaaaga
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aagaaataa
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           20
Ile Ile Lys Lys Leu Lys Leu Asp Pro Glu Ala Arg Ala Ser Glu Leu
                           40
Thr Glu Glu Glu Val Gly Arg Leu Asn Ser Leu Leu Gln Ser Glu Tyr
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Thr Val Glu Gly Asp Leu Arg Arg Arg Val Gln Ser Asp Ile Lys Arg
                   70
                                        75
Leu Ile Ala Ile His Ser Tyr Arg Gly Gln Arg His Arg Leu Ser Leu
               85
                                    90
Pro Val Arg Gly Gln Arg Thr Lys Thr Asn Ser Arg Thr Arg Lys Gly
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Lys Arg Lys Thr Val Ala Gly Lys Lys
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Leu Phe Val Asn Lys Met Leu Ala Gln Pro Phe Leu Ser Ser Gln Thr
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Lys Ala Asn Met Gly
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                                                                       120
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                                                   30
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Leu Phe Val Asn Lys Met Leu Ala Lys Pro Phe Leu Ser Ser Gln Thr
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                           40
Lys Ala Asn Met Gly
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      <211> 33
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      <211> 53
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       <211> 30
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1
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cgccgtgggc gatttagcga aaaatgattc ttctattcaa gtacgcatca ctgcttatcg 180
tgctgcagcc gtgttggaga tacaagatct tgtgcctcat ttacgagttg tagtccaaaa 240
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      <220>
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qttqttctaq ctttqqtacq aqaaqqtqat tctaaqccct acqcqattaq ttatqqatac 420
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aaaagetetg ggagcatgtt ettagtetea geagatatta ttgeateaag aatggaagge 240
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gaggtaatac ctcaaacaaa cgcttaaaca atttttattg gatttttctt ataggtttta 540
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17

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19

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WO 02/08267 PC

26

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Ala Gly Gly Ile Glu Gly Thr Glu Tyr Pro Leu Leu Ala Asp Pro Ser 100 105 110

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WO 02/08267

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PCT/US01/23121 WO 02/08267 37

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 Val Gln Ser Ala Gln Ser Phe Phe Ser Tyr Met Lys Ala Ala Ser Gln
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WO 02/08267 PCT/US01/23121

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WO 02/08267

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WO 02/08267

43

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660

720

780 840

897

44

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His Lys Arg Arg Ala Ala Ala Val Cys Ser Phe Ile Gly Gly Ile
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Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn
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gegggetett eegeacacat tacagettee caagtgteea aaggattagg ggatgegaga actgtteteg etttagggaa tgeetttaae ggagegttge eaggaacagt teaaagtgeg etegtageag atetttgtgt gteetataag egeagagegg etgeggetgt etgtagette ateggaggaa ttacetacet egegacatte ggagetatee gteegattet gtettettee eaaatgetgg egeaacegtt tettettee eaaactaaag eaaattgtegg eaggaggaaaa eegeeggetge etgtggtgg gttetggaet egetaeagt geggaaagag eagattgega egeeggetge etgtggtgg gttetggaet egetaeagt egeggaaagag eagattgega egeeggetge etgtggtgg gttetggaet egetaeagt egeggaaatgtegg gagaggaaaa tgeetggaggagaagagggggaggaggaggaggaggaggaggag															
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Met		400>		Cys	G) v	Ara	T.O.I.	G) v	Sar	G1v	ሞኮሎ	Clu	λen	71.	Ton
1	NT.	Ara	116	5	Gry	Arg	Бец	GTĀ	10	дту	1111	сту	ASII	15	пеп
Lys	Ala	Phe	Phe 20	Thr	Gln	Pro	Ser	Asn 25	Lys	Met	Ala	Arg	Val 30	Val	Asn
Lys	Thr	Lys 35		Met	Asp	Lys	Thr 40		Lys	Val	Ala	Lys 45		Ala	Ala
Glu	Leu 50	Thr	Ala	Asn	Ile	Leu 55	Glu	Gln	Ala	Gly		Ala	Gly	Ser	Ser
Ala 65		Ile	Thr	Ala	Ser 70		Val	Ser	Lys	Gly 75	60 Leu	Gly	Asp	Ala	Arg 80
Thr	۷al	Leu	Ala	Leu 85	Gly	Asn	Ala	Phe	Asn 90	Gly	Ala	Leu	Pro	Gly 95	
Val	Gln	Ser	Ala 100	Gln	Ser	Phe	Phe	Ser 105		Met	Lys	Ala	Ala 110		Gln
Lys	Pro	Gln 115	Glu	Gly	Asp	Glu	Gly 120		Val	Ala	Asp	Leu 125		Val	Ser
His	Lys 130		Arg	Ala	Ala	Ala 135		Val	Cys	Ser	Phe 140		Gly	Gly	Ile
Thr 145	Tyr	Leu	Ala	Thr	Phe 150		Ala	Ile	Arg	Pro 155		Leu	Phe	Val	Asn 160
	Met	Leu	Ala	Gln		Phe	Leu	Ser			Thṛ	Lys	Ala	Asn	
Gly	Ser	Ser	Val 180	165 Ser	Tyr	11e	Met	Ala 185	170 Ala	Asn	His	Ala		175 Phe	Val
Val	Gly			Leu	Ala	Ile			Glu	Arg	Ala		190 Cys	Glu	Ala
Arg	Cys 210	195 Ala	Arg	Ile	Ala		200 Glu	Glu	Ser	Ser		205 Glu	Leu	Ser	Gly
		Asn	Ala	Суз		215 Arg	Arg	Val	Ala		220 Glu	Lys	Ala	Lys	
225 Phe	Thr	Arg	Ile	Lys	230 Tyr	Ala	Leu	Leu	Thr	235 Met	Leu	Glu	Lys	Phe	240 Leu
				245 Asp					250					255	
			260					265					270		
		275		Ile			280			ınr	rne	Thr 285	Ser	Ala	va⊥
Ile	Gly 290	Leu	Trp	Thr	Phe	Cys 295	Asn	Arg	Val			,			
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<211> 897 <212> DNA

<213> Chlamydia

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<211> 298

<212> PRT

<213> Chlamydia

<400> 135

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                                                                         120
ataaaggttg ggaagtctgc tgctgaatta acggcgagta ttttagagca aactgggggg
                                                                         180
gcagggactg atgcacatgt tacggcggcc aaggtgtcta aagcacttgg ggacgcgcga
                                                                         240
acagtaatgg ctctagggaa tgtcttcaat gggtctgtgc cagcaaccat tcaaagtgcg
                                                                         300
cgaagctgtc tcgcccattt acgagcggcc ggcaaagaag aagaaacatg ctccaaggtg
                                                                         360
aaagatetet gtgtttetea tagaegaaga getgeggetg aggettgtaa tgttattgga
                                                                         420
ggagcaactt atattacaac tttcggagcg attcgtccga cattactcgt taacaagctt
                                                                         480
cttgccaaac cattcctttc ctcccaagcc aaagaagggt tgggagcttc tgttggttat
                                                                         540
atcatggcag cgaaccatgc ggcatctgtg ctttgggtctg ctttaagtat tagcgcagaa
                                                                         600
agagcagact gtgaagagcg gtgtgatcgc attcgatgta gtgaggatgg tgaaatttgc gaaggcaata aattaacagc tatttcggaa gagaaggcta gatcatggac tctcattaag
                                                                         660
                                                                         720
tacagattee ttactatgat agaaaaacta tttgagatgg tggeggatat etteaagtta
                                                                         780
attectttge caatttegea tggaattegt getattgttg etgegggatg taegttgaet
                                                                         840
tctgcagtta ttggcttagg tactttttgg tctagagcat aa
                                                                         882
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Ser Ala Lys Gly Leu Asp Arg Ser Ile Lys Val Gly Lys Ser Ala Ala 35 40 45
Glu Leu Thr Ala Ser Ile Leu Glu Gln Thr Gly Gly Ala Gly Thr Asp
                       55
Ala His Val Thr Ala Ala Lys Val Ser Lys Ala Leu Gly Asp Ala Arg
                    70
                                         75
Thr Val Met Ala Leu Gly Asn Val Phe Asn Gly Ser Val Pro Ala Thr
                                     90
Ile Gln Ser Ala Arg Ser Cys Leu Ala His Leu Arg Ala Ala Gly Lys
                                 105
            100
                                                     110
Glu Glu Glu Thr Cys Ser Lys Val Lys Asp Leu Cys Val Ser His Arg
                           120
       115
                                                 125
Arg Arg Ala Ala Ala Glu Ala Cys Asn Val Ile Gly Gly Ala Thr Tyr
                       135
Ile Thr Thr Phe Gly Ala Ile Arg Pro Thr Leu Leu Val Asn Lys Leu
                    150
                                         155
Leu Ala Lys Pro Phe Leu Ser Ser Gln Ala Lys Glu Gly Leu Gly Ala
                165
                                                          175
                                     170
Ser Val Gly Tyr Ile Met Ala Ala Asn His Ala Ala Ser Val Leu Gly
            180
                                185
Ser Ala Leu Ser Ile Ser Ala Glu Arg Ala Asp Cys Glu Glu Arg Cys
                            200
                                                 205
Asp Arg Ile Arg Cys Ser Glu Asp Gly Glu Ile Cys Glu Gly Asn Lys
                       215
                                            220
Leu Thr Ala Ile Ser Glu Glu Lys Ala Arg Ser Trp Thr Leu Ile Lys
                    230
                                         235
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Tyr Arg Phe Leu Thr Met Ile Glu Lys Leu Phe Glu Met Val Ala Asp
         245
                           250
Ile Phe Lys Leu Ile Pro Leu Pro Ile Ser His Gly Ile Arg Ala Ile
                           265
Val Ala Ala Gly Cys Thr Leu Thr Ser Ala Val Ile Gly Leu Gly Thr
Phe Trp Ser Arg Ala
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Arg Ala Ala Ala Val Cys Ser Phe Ile Gly Gly Ile Thr Tyr Leu
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Arg Pro
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Met Leu
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Ser Gln
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Ser
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Ser Phe Ile Gly Gly Ile Thr Tyr Leu
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Cys Ser Phe Ile Gly Gly Ile Thr
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1 10
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Gly Phe Ile Gly Gly Ile Thr Tyr Leu
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PCT/US01/23121 WO 02/08267

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Ser Val Ala Ser
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     <211> 20
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1
Thr Ser Arg His
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Arg Phe Cys Leu
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      <211> 20
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Arg Asn Arg Phe
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· Gln Ile Trp Asp
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       <211> 53
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                                   10
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                        25
 Phe Cys Leu Ser Thr Lys Cys Trp Arg Asn Arg Phe Phe Leu Pro Lys
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 Leu Lys Gln Ile Trp
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 Ile Gly Gly Ile Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile
         20
                             25
                                            30
 Leu Phe Val Asn Lys Met Leu Ala Gln Pro Phe Leu Ser Ser Gln Ile
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                  40
 Lys Ala Asn Met
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ser Phe lie Gly Gly 1.	re Tit lât men

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                                                                          120
aatcatgtcg tctgtacatt ttttgaggac tgtaccatgg agagcctctt tcctgctctt
                                                                          180
tgtgctcatg catcacaaga cgatcctttg tatgtacttg gaaattccta ctgttgqttc
                                                                          240
gtatctaaac tccatatcac ggaccccaaa gaggctcttt ttaaagaaaa aggagatctt
                                                                          300
tocattoaaa actttogett cettteette acagattget ettecaagga aageteteet
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tctattattc atcaaaagaa tggtcagtta tccttgcgca ataatggtag catgagtttc
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                                                                         1260
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                                                                         1440
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                                                                         1500
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ettactetee ctaaagagea ateteatita catetteetg atgggaacet etetteteae
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                                                                         1740
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                                                                         2040
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                                                                         2100
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                                                                         2160
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                                                                         2220
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                                                                         2280
gacggttttg aggagagttc gggagagatt cggtcctttt ctgccagctc tttcagaaat
                                                                         2340
atticactic ctataggaat aacattigaa aaaaaatccc aaaaaacacg aacctactat
                                                                         2400
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                                                                         2460
ttactcaaaa atgccgtctc ctgggatgct cctatggcga acttggattc acgagcctac
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gttttaggga gaggacactc					240
gctctaagta atagcgctgc	tgatggactg	tttactattg	agggttttaa	agaattatcc	300
ttttccaatt gcaattcatt					360
cagactccga cgacaacatc					420
ttgttactca ataatgagaa					480 540
gctatagatg ctaagagctt aatactgctc aagctgatgg					600
aacgaggete ctattgeett					660
gctgttcagg atgggcagca					720
agtttttcca gaaatactgc					780
atttactcct acgggaacgt					840
gttgcttctc ctgtttacat		•		_	900
agtaataatt acggagatgg aataactctg gatcagtttc					960 1020
gctgctggga aagggggagc					1020
gtacaatttt taaggaatat					1140
gageteagtt tatetgetga					1200
gccaaagaga atgctgccga					1260
ggatcgggag ggaaaataac					1320
gateceateg agatggeaaa					1380
attaacgatg gtgaaggata taccaaaatg ttacgataga					1440 1500
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tttgtaactc cacaaccacc					1620
aatctgcatt tgtctctttc					1680
aatcctccag cgcaagattc					1740
acaattagtg ggcctatctt					1800
tggctaggtt ctaatcaaaa					1860
gctaatgccc catcagattt agctggaagc ttgcgtggga					1920 1980
acatggacta aaactgggta					2040
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gatgggcgct cttattgtcg					2160
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                                                                         4740
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                                                                         4800
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ccttctatct acagaaataa tcctgtctgt aaatatcggg tattgtcttc gaatgaagct
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ctatatcttg gtcccttctg gactctctac ggaaactata ctatcgatgt aggcatgtat
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           20
                                                    30
Thr Ala Leu Leu Thr Lys Asn Pro Asn His Val Val Cys Thr Phe Phe
                           40
                                                45
Glu Asp Cys Thr Met Glu Ser Leu Phe Pro Ala Leu Cys Ala His Ala
                        55
Ser Gln Asp Asp Pro Leu Tyr Val Leu Gly Asn Ser Tyr Cys Trp Phe
                                        75
Val Ser Lys Leu His Ile Thr Asp Pro Lys Glu Ala Leu Phe Lys Glu
                                    90
                85
Lys Gly Asp Leu Ser Ile Gln Asn Phe Arg Phe Leu Ser Phe Thr Asp
                                105
            100
Cys Ser Ser Lys Glu Ser Ser Pro Ser Ile Ile His Gln Lys Asn Gly
                           120
                                                125
Gln Leu Ser Leu Arg Asn Asn Gly Ser Met Ser Phe Cys Arg Asn His
                        135
                                            140
Ala Glu Gly Ser Gly Gly Ala Ile Ser Ala Asp Ala Phe Ser Leu Gln
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Gl	у (Gly	Ala	Ile 180		Ala	Gln	Thr	Phe 185		Leu	Ser	Arg	Asn 190	Val	Ser
Pr	: o	Ile	Ser 195	Phe	Ala	Arg	Asn	Arg 200	Ala	Asp	Leu	Asn	Gly 205	Gly	Ala	Ile
СУ		Cys 210			Leu	Ile	Cys 215	Ser	Gly	Asn	Val	Asn 220	Pro	Leu	Phe	Phe
22	25	_				230					235			Ile		240
					245					250				Asn	255	
				260					265					Ala 270		
		-	275					280					285	Phe		
		290					295					300		Gly		
30)5					310					315			Asn		320
		_			325					330				Tyr	335	
_		-		340					345					Asp 350		
			355					360					365	Ser		
		370					375					380		Ala		
38	35					390					395			Ile		400
					405					410				Leu	415	
				420					425					Val 430		
	_	_	435					440					445	Pro		
		450					455					460		Ser		
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					485					490				Ile	495	
				500					505					Leu 510		
			515					520					525	Glu		
		530					535					540		Gly		
5	45					550					555			His	•	560
					565					570				Glu	575	
				580					585					Asp 590		
			595					600			•		605			
		610					615					620		Val		
6	25					630					635			Leu		640
L	eu	Phe	Gly	Ile	Ser	Thr	His	Ser	Leu	Asp	Asp	His	Ser	Phe	Cys	Leu

63

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Ala Ala Gly Gln Leu Leu Gly Lys Ser Ser Asp Ser Phe Ile Thr Ser
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Thr Glu Thr Thr Ser Tyr Ile Ala Thr Val Gln Ala Gln Leu Ala Thr
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Ser Leu Met Lys Ile Ser Ala Gln Ala Cys Tyr Asn Glu Ser Ile His
690 695 700
Glu Leu Lys Thr Lys Tyr Arg Ser Phe Ser Lys Glu Gly Phe Gly Ser 705 710 715 720
Trp His Ser Val Ala Val Ser Gly Glu Val Cys Ala Ser Ile Pro Ile
                                 730
Val Ser Asn Gly Ser Gly Leu Phe Ser Ser Phe Ser Ile Phe Ser Lys
          740
                              745
Leu Gln Gly Phe Ser Gly Thr Gln Asp Gly Phe Glu Glu Ser Ser Gly
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Glu Ile Arg Ser Phe Ser Ala Ser Ser Phe Arg Asn Ile Ser Leu Pro 770 775 780
Ile Gly Ile Thr Phe Glu Lys Lys Ser Gln Lys Thr Arg Thr Tyr Tyr 785 790 795 800
                                    795
Tyr Phe Leu Gly Ala Tyr Ile Gln Asp Leu Lys Arg Asp Val Glu Ser
                                 810
Gly Pro Val Val Leu Leu Lys Asn Ala Val Ser Trp Asp Ala Pro Met
          820 825
                                              830
Ala Asn Leu Asp Ser Arg Ala Tyr Met Phe Arg Leu Thr Asn Gln Arg
     835 . 840
                                          845
Ala Leu His Arg Leu Gln Thr Leu Leu Asn Val Ser Cys Val Leu Arg
 850 855
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Pro Tyr Thr Val Ile Gly Asp Pro Ser Gly Thr Thr Val Phe Ser Ala
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           20
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                                             45
                          40
Leu Ser Cys Phe Gly Asn Leu Leu Gly Ser Phe Thr Val Leu Gly Arg
                    55
Gly His Ser Leu Thr Phe Glu Asn Ile Arg Thr Ser Thr Asn Gly Ala
Ala Leu Ser Asn Ser Ala Ala Asp Gly Leu Phe Thr Ile Glu Gly Phe
                               90
            85
Lys Glu Leu Ser Phe Ser Asn Cys Asn Ser Leu Leu Ala Val Leu Pro
                             105
Ala Ala Thr Thr Asn Lys Gly Ser Gln Thr Pro Thr Thr Thr Ser Thr
    115
                          120
                                             125
Pro Ser Asn Gly Thr Ile Tyr Ser Lys Thr Asp Leu Leu Leu Asn 130 135
Asn Glu Lys Phe Ser Phe Tyr Ser Asn Leu Val Ser Gly Asp Gly Gly
                                     155
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Ala Ile Asp Ala Lys Ser Leu Thr Val Gln Gly Ile Ser Lys Leu Cys 170 165 Val Phe Gln Glu Asn Thr Ala Gln Ala Asp Gly Gly Ala Cys Gln Val 180 185 Val Thr Ser Phe Ser Ala Met Ala Asn Glu Ala Pro Ile Ala Phe Val 200 205 195 Ala Asn Val Ala Gly Val Arg Gly Gly Gly Ile Ala Ala Val Gln Asp 215 220 Gly Gln Gln Gly Val Ser Ser Ser Thr Ser Thr Glu Asp Pro Val Val 230 235 Ser Phe Ser Arg Asn Thr Ala Val Glu Phe Asp Gly Asn Val Ala Arg 245 250 Val Gly Gly Ile Tyr Ser Tyr Gly Asn Val Ala Phe Leu Asn Asn 260 265 270 265 Gly Lys Thr Leu Phe Leu Asn Asn Val Ala Ser Pro Val Tyr Ile Ala 285 280 Ala Lys Gln Pro Thr Ser Gly Gln Ala Ser Asn Thr Ser Asn Asn Tyr 290 295 300 Gly Asp Gly Gly Ala Ile Phe Cys Lys Asn Gly Ala Gln Ala Gly Ser 310 315 Asn Asn Ser Gly Ser Val Ser Phe Asp Gly Glu Gly Val Val Phe Phe 325 330 335 330 325 Ser Ser Asn Val Ala Ala Gly Lys Gly Gly Ala Ile Tyr Ala Lys Lys 340 345 350340 Leu Ser Val Ala Asn Cys Gly Pro Val Gln Phe Leu Arg Asn Ile Ala 360 Asn Asp Gly Gly Ala Ile Tyr Leu Gly Glu Ser Gly Glu Leu Ser Leu 370 375 380 Ser Ala Asp Tyr Gly Asp Ile Ile Phe Asp Gly Asn Leu Lys Arg Thr 385 390 395 400 Ala Lys Glu Asn Ala Ala Asp Val Asn Gly Val Thr Val Ser Ser Gln 405 410 Ala Ile Ser Met Gly Ser Gly Gly Lys Ile Thr Thr Leu Arg Ala Lys 425 420 Ala Gly His Gln Ile Leu Phe Asn Asp Pro Ile Glu Met Ala Asn Gly 440 445 435 Asn Asn Gln Pro Ala Gln Ser Ser Lys Leu Leu Lys Ile Asn Asp Gly 450 455 460 Glu Gly Tyr Thr Gly Asp Ile Val Phe Ala Asn Gly Ser Ser Thr Leu 465 470 475 480 470 475 Tyr Gln Asn Val Thr Ile Glu Gln Gly Arg Ile Val Leu Arg Glu Lys 490 Ala Lys Leu Ser Val Asn Ser Leu Ser Gln Thr Gly Gly Ser Leu Tyr 500 505 Met Glu Ala Gly Ser Thr Leu Asp Phe Val Thr Pro Gln Pro Pro Gln 525 520 Gln Pro Pro Ala Ala Asn Gln Leu Ile Thr Leu Ser Asn Leu His Leu 530 535 540 Ser Leu Ser Ser Leu Leu Ala Asn Asn Ala Val Thr Asn Pro Pro Thr 550 555 Asn Pro Pro Ala Gln Asp Ser His Pro Ala Val Ile Gly Ser Thr Thr 570 565 Ala Gly Ser Val Thr Ile Ser Gly Pro Ile Phe Phe Glu Asp Leu Asp 590 585 580 Asp Thr Ala Tyr Asp Arg Tyr Asp Trp Leu Gly Ser Asn Gln Lys Ile 600 595 Asn Val Leu Lys Leu Gln Leu Gly Thr Lys Pro Pro Ala Asn Ala Pro 615 620 Ser Asp Leu Thr Leu Gly Asn Glu Met Pro Lys Tyr Gly Tyr Gln Gly 625 630 635 640 630 Ser Trp Lys Leu Ala Trp Asp Pro Asn Thr Ala Asn Asn Gly Pro Tyr

655 645 650 Thr Leu Lys Ala Thr Trp Thr Lys Thr Gly Tyr Asn Pro Gly Pro Glu 660 665 670 665 Arg Val Ala Ser Leu Val Pro Asn Ser Leu Trp Gly Ser Ile Leu Asp 680 685 Ile Arg Ser Ala His Ser Ala Ile Gln Ala Ser Val Asp Gly Arg Ser 690 695 700 Tyr Cys Arg Gly Leu Trp Val Ser Gly Val Ser Asn Phe Phe Tyr His 710 715 Asp Arg Asp Ala Leu Gly Gln Gly Tyr Arg Tyr Ile Ser Gly Gly Tyr 725 730 Ser Leu Gly Ala Asn Ser Tyr Phe Gly Ser Ser Met Phe Gly Leu Ala 740 745 750 Phe Thr Glu Val Phe Gly Arg Ser Lys Asp Tyr Val Val Cys Arg Ser 765 755 760 Asn His His Ala Cys Ile Gly Ser Val Tyr Leu Ser Thr Gln Gln Ala 770 780 775 780 Leu Cys Gly Ser Tyr Leu Phe Gly Asp Ala Phe Ile Arg Ala Ser Tyr 790 795 Gly Phe Gly Asn Gln His Met Lys Thr Ser Tyr Thr Phe Ala Glu Glu 805 810 Ser Asp Val Arg Trp Asp Asn Asn Cys Leu Ala Gly Glu Ile Gly Ala 825 820 830 Gly Leu Pro Ile Val Ile Thr Pro Ser Lys Leu Tyr Leu Asn Glu Leu 840 Arg Pro Phe Val Gln Ala Glu Phe Ser Tyr Ala Asp His Glu Ser Phe 855 860 Thr Glu Glu Gly Asp Gln Ala Arg Ala Phe Lys Ser Gly His Leu Leu 870 875 Asn Leu Ser Val Pro Val Gly Val Lys Phe Asp Arg Cys Ser Ser Thr 885 890 His Pro Asn Lys Tyr Ser Phe Met Ala Ala Tyr Ile Cys Asp Ala Tyr 905 Arg Thr Ile Ser Gly Thr Glu Thr Thr Leu Leu Ser His Gln Glu Thr 915 920 925 Trp Thr Thr Asp Ala Phe His Leu Ala Arg His Gly Val Val Val Arg 930 935 940 Gly Ser Met Tyr Ala Ser Leu Thr Ser Asn Ile Glu Val Tyr Gly His 945 950 955 960 Gly Arg Tyr Glu Tyr Arg Asp Ala Ser Arg Gly Tyr Gly Leu Ser Ala 965 . 970 Gly Ser Lys Val Xaa Phe 980 <210> 177 <211> 964 <212> PRT <213> Chlamydia <400> 177 Met Lys Lys Ala Phe Phe Phe Leu Ile Gly Asn Ser Leu Ser Gly 5 10 Leu Ala Arg Glu Val Pro Ser Arg Ile Phe Leu Met Pro Asn Ser Val 20 30 Pro Asp Pro Thr Lys Glu Ser Leu Ser Asn Lys Ile Ser Leu Thr Gly 40 35 Asp Thr His Asn Leu Thr Asn Cys Tyr Leu Asp Asn Leu Arg Tyr Ile 55 60 Leu Ala Ile Leu Gln Lys Thr Pro Asn Glu Gly Ala Ala Val Thr Ile 70

Thr Asp Tyr Leu Ser Phe Phe Asp Thr Gln Lys Glu Gly Ile Tyr Phe

				85					90					95	
Ala	Lys	Asn	Leu 100	Thr	Pro	Glu	Ser	Gly 105	Gly	Ala	Ile	Gly	Tyr 110	Ala	Ser
Pro	Asn	Ser 115	Pro	Thr	Val	Glu	Ile 120	Arg	Asp	Thr	Ile	Gly 125	Pro	Val	Ile
Phe	Glu 130	Asn	Asn	Thr	Суѕ	Cys 135	Arg	Leu	Phe	Thr	Trp 140	Arg	Asn	Pro	Tyr
Ala 145	Ala	Asp	Lys	Ile	Arg 150	Glu	Gly	Gly	Ala	Ile 155	His	Ala	Gln	Asn	Leu 160
Tyr	Ile	Asn	His	Asn 165	His	Asp	Val		Gly 170	Phe	Met	Lys	Asn	Phe 175	Ser
Tyr	Val	Gln	Gly 180	Gly	Ala	Ile	Ser	Thr 185	Ala	Asn	Thr	Phe	Val 190	Val	Ser
Glu	Asn	Gln 195	Ser	Суѕ	Phe	Leu	Phe 200	Met	Asp	Asn	Ile	Cys 205	Ile	Gln	Thr
Asn	Thr 210	Ala	Gly	Lys	Gly	Gly 215	Ala	Ile	Tyr	Ala	Gly 220	Thr	Ser	Asn	Ser
Phe 225	Glu	Ser	Asn	Asn	Cys 230	Asp	Leu	Phe	Phe	Ile 235	Asn	Asn	Ala	Суѕ	Cys 240
Ala	Gly	Gly	Ala	Ile 245	Phe	Ser	Pro	Ile	Cys 250	Ser	Leu	Thr	Gly	Asn 255	Arg
			260		Tyr			265					270		
Ala	Ser	Ser 275	Glu	Ala	Ser	Asp	Gly 280	Gly	Ala	Ile	Lys	Val 285	Thr,	Thr	Arg
	290				Asn	295	-	-			300		-		
Thr 305	ГÀЗ	Asn	Tyr	Gly	Gly 310	Ala	Ile	Tyr	Ala	Pro 315	Val	Val	Thr	Leu	Val 320
_		_		325	Tyr				330					335	•
			340		Asp			345			_		350		_
		355			Phe		360					365			
	370				Thr	375					380				
385					Ser 390					395			_		400
				405	Tyr -				410					415	
			420		Lys			425			_		430		
		435			Asn		440					445			
	450				Pro	455					460				
465					Leu 470					475				_	480
				485	Asn				490					495	
			500		Ser			505					510		_
		515			Ile		520					525			
	530				Asn	535					540		_		
545					Ser 550					555			_	_	560
Gly	Asn	Ser	Pro	Tyr 565	Glu	Ser	Thr	Asp	Leu 570	Thr	His	Ala	Leu	Ser 575	Ser

Gln Pro Met Leu Ser Ile Ser Glu Ala Ser Asp Asn Gln Leu Gln Ser 580 585 Glu Asn·Ile Asp Phe Ser Gly Leu Asn Val Pro His Tyr Gly Trp Gln 600 605 Gly Leu Trp Thr Trp Gly Trp Ala Lys Thr Gln Asp Pro Glu Pro Ala 610 620 Ser Ser Ala Thr Ile Thr Asp Pro Gln Lys Ala Asn Arg Phe His Arg 625 630 635 625 630 635 Thr Leu Leu Thr Trp Leu Pro Ala Gly Tyr Val Pro Ser Pro Lys 650 655 His Arg Ser Pro Leu Ile Ala Asn Thr Leu Trp Gly Asn Met Leu Leu. 660 665 670 Ala Thr Glu Ser Leu Lys Asn Ser Ala Glu Leu Thr Pro Ser Gly His 680 675 685 Pro Phe Trp Gly Ile Thr Gly Gly Gly Leu Gly Met Met Val Tyr Gln 695 Asp Pro Arg Glu Asn His Pro Gly Phe His Met Arg Ser Ser Gly Tyr 705 710 715 720 Ser Ala Gly Met Ile Ala Gly Gln Thr His Thr Phe Ser Leu Lys Phe $725 \hspace{1.5cm} 730 \hspace{1.5cm} 735$ Ser Gln Thr Tyr Thr Lys Leu Asn Glu Arg Tyr Ala Lys Asn Asn Val 740 745 750 740 745 Ser Ser Lys Asn Tyr Ser Cys Gln Gly Glu Met Leu Phe Ser Leu Gln 760 Glu Gly Phe Leu Leu Thr Lys Leu Val Gly Leu Tyr Ser Tyr Gly Asp
770 780 His Asn Cys His His Phe Tyr Thr Gln Gly Glu Asn Leu Thr Ser Gln 785 790 795 800 Gly Thr Phe Arg Ser Gln Thr Met Gly Gly Ala Val Phe Phe Asp Leu 805 810 815 Pro Met Lys Pro Phe Gly Ser Thr His Ile Leu Thr Ala Pro Phe Leu 825 Gly Ala Leu Gly Ile Tyr Ser Ser Leu Ser His Phe Thr Glu Val Gly 835 840 845 Ala Tyr Pro Arg Ser Phe Ser Thr Lys Thr Pro Leu Ile Asn Val Leu 850 855 860 Val Pro Ile Gly Val Lys Gly Ser Phe Met Asn Ala Thr His Arg Pro 865 870 875 880 Gln Ala Trp Thr Val Glu Leu Ala Tyr Gln Pro Val Leu Tyr Arg Gln 885 890 895 Glu Pro Gly Ile Ala Thr Gln Leu Leu Ala Ser Lys Gly Ile Trp Phe 900 905 910 Gly Ser Gly Ser Pro Ser Ser Arg His Ala Met Ser Tyr Lys Ile Ser 915 920 925 Gln Gln Thr Gln Pro Leu Ser Trp Leu Thr Leu His Phe Gln Tyr His 930 935 940 Gly Phe Tyr Ser Ser Ser Thr Phe Cys Asn Tyr Leu Asn Gly Glu Ile 950 Ala Leu Arg Phe

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<211> 1530

<212> PRT

<213> Chlamydia

<400> 178

Met Ser Ser Glu Lys Asp Ile Lys Ser Thr Cys Ser Lys Phe Ser Leu

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20 25 30

Val Asp Leu His Ala Gly Gly Gln Ser Val Asn Glu Leu Val Tyr Val Gly Pro Gln Ala Val Leu Leu Asp Gln Ile Arg Asp Leu Phe Val Gly Ser Lys Asp Ser Gln Ala Glu Gly Gln Tyr Arg Leu Ile Val Gly Asp Pro Ser Ser Phe Gln Glu Lys Asp Ala Asp Thr Leu Pro Gly Lys Val Glu Gln Ser Thr Leu Phe Ser Val Thr Asn Pro Val Val Phe Gln Gly Val Asp Gln Gln Asp Gln Val Ser Ser Gln Gly Leu Ile Cys Ser Phe Thr Ser Ser Asn Leu Asp Ser Pro Arg Asp Gly Glu Ser Phe Leu 130 140 Gly Ile Ala Phe Val Gly Asp Ser Ser Lys Ala Gly Ile Thr Leu Thr Asp Val Lys Ala Ser Leu Ser Gly Ala Ala Leu Tyr Ser Thr Glu Asp Leu Ile Phe Glu Lys Ile Lys Gly Gly Leu Glu Phe Ala Ser Cys Ser Ser Leu Glu Gln Gly Gly Ala Cys Ala Ala Gln Ser Ile Leu Ile His 195 200 205Asp Cys Gln Gly Leu Gln Val Lys His Cys Thr Thr Ala Val Asn Ala Glu Gly Ser Ser Ala Asn Asp His Leu Gly Phe Gly Gly Gly Ala Phe Phe Val Thr Gly Ser Leu Ser Gly Glu Lys Ser Leu Tyr Met Pro Ala Gly Asp Met Val Val Ala Asn Cys Asp Gly Ala Ile Ser Phe Glu Gly Asn Ser Ala Asn Phe Ala Asn Gly Gly Ala Ile Ala Ala Ser Gly Lys Val Leu Phe Val Ala Asn Asp Lys Lys Thr Ser Phe Ile Glu Asn Arg Ala Leu Ser Gly Gly Ala Ile Ala Ala Ser Ser Asp Ile Ala Phe Gln Asn Cys Ala Glu Leu Val Phe Lys Gly Asn Cys Ala Ile Gly Thr Glu Asp Lys Gly Ser Leu Gly Gly Gly Ala Ile Ser Ser Leu Gly Thr Val Leu Leu Gln Gly Asn His Gly Ile Thr Cys Asp Lys Asn Glu Ser Ala Ser Gln Gly Gly Ala Ile Phe Gly Lys Asn Cys Gln Ile Ser Asp Asn Glu Gly Pro Val Val Phe Arg Asp Ser Thr Ala Cys Leu Gly Gly Ala Ile Ala Ala Gln Glu Ile Val Ser Ile Gln Asn Asn Gln Ala Gly Ile Ser Phe Glu Gly Gly Lys Ala Ser Phe Gly Gly Ile Ala Cys Gly Ser Phe Ser Ser Ala Gly Gly Ala Ser Val Leu Gly Thr Ile Asp Ile Ser Lys Asn Leu Gly Ala Ile Ser Phe Ser Arg Thr Leu Cys Thr Thr Ser Asp Leu Gly Gln Met Glu Tyr Gln Gly Gly Gly Ala Leu Phe Gly Glu Asn Ile Ser Leu Ser Glu Asn Ala Gly Val Leu Thr Phe Lys Asp Asn Ile Val Lys Thr Phe Ala Ser Asn Gly Lys Ile Leu Gly Gly Gly Ala Ile Leu Ala Thr Gly Lys Val Glu Ile Thr Asn Asn Ser Gly

		515					520					525			
Gly	Ile 530	Ser	Phe	Thr	Gly	Asn 535	Ala	Arg	Ala	Pro	Gln 540	Ala	Leu	Pro	Thr
Gln 545	Glu	Glų	Phe	Pro	Leu 550	Phe	Ser	Lys	Lys	Glu 555		Arg	Pro	Leu	Ser 560
Ser	Gly	Tyr	Ser	Gly 565		Gly	Ala	Ile	Leu 570		Arg	Glu	Val	Ala 575	
Leu	His	Asn	Ala 580	Ala	Val	Val	Phe	Glu 585	Gln	Asn	Arg	Leu	Gln 590		Ser
Glu	Glu	Glu 595		Thr	Leu	Leu	Gly 600		Cys	Gly	Gly	Gly 605		Val	His
Gly	Met 610	Asp	Ser	Thr	Ser	Ile 615	Val	Gly	Asn	Ser	Ser 620	Val	Arg	Phe	Gly
Asn 625	Asn	Tyr	Ala	Met	Gly 630	Gln	Gly	Val	Ser	Gly 635	Gly	Ala	Leu	Leu	Ser 640
Lys	Thr	Val	Gln	Leu 645	Ala	Gly	Asn	Gly	Ser 650	Val	Asp	Phe	Ser	Arg 655	Asn
Ile	Ala	Ser	Leu 660	Gly	Gly	Gly	Ala	Leu 665	Gln	Ala	Ser	Glu	Gly 670	Asn	Cys
Glu	Leu	Val 675	Asp	Asn	Gly	Tyr	Val 680	Leu	Phe	Arg	Asp	Asn 685	Arg	Gly	Arg
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705					710				Asp	715				_	720
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			740					745	Phe				750		
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	770					775			Ser		780				
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				805					Asn 810					815	-
			820					825	Arg				830		-
-		835					840		Gly			845	-		
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865	_			_	870				Thr	875					880
				885					Cys 890		_			895	
			900					905	Glu				910		
		915					920		Ala		_	925	_		
	930					935			Thr		940				
945					950	•			Leu	955					960
				965					Ile 970			_		975	
_			980					985	Glu				990		
Gln	Cys	11e 995	His	Val	Gln	Gln	Gly 1000		Leu	Glu	Leu	Leu 1005		Gly	Ala

Thr Leu Cys Ser Tyr Gly Phe Lys Gln Asp Ala Gly Ala Lys Leu Val 1010 1015 1020 Leu Ala Ala Gly Ser Lys Leu Lys Ile Leu Asp Ser Gly Thr Pro Val 1030 1035 Gln Gly His Ala Ile Ser Lys Pro Glu Ala Glu Ile Glu Ser Ser Ser 1045 1050 1055 Glu Pro Glu Gly Ala His Ser Leu Trp Ile Ala Lys Asn Ala Gln Thr 1060 1065 1070 Thr Val Pro Met Val Asp Ile His Thr Ile Ser Val Asp Leu Ala Ser 1075 1080 1085 Phe Ser Ser Gln Gln Glu Gly Thr Val Glu Ala Pro Gln Val Ile 1090 1095 1100 Val Pro Gly Gly Ser Tyr Val Arg Ser Gly Glu Leu Asn Leu Glu Leu 1105 1110 1115 1120 Val Asn Thr Thr Gly Thr Gly Tyr Glu Asn His Ala Leu Leu Lys Asn 1125 1130 1135 Glu Ala Lys Val Pro Leu Met Ser Phe Val Ala Ser Ser Asp Glu Ala 1140 1145 1150 Ser Ala Glu Ile Ser Asn Leu Ser Val Ser Asp Leu Gln Ile His Val 1155 1160 1165 Ala Thr Pro Glu Ile Glu Glu Asp Thr Tyr Gly His Met Gly Asp Trp 1170 1175 1180 Ser Glu Ala Lys Ile Gln Asp Gly Thr Leu Val Ile Asn Trp Asn Pro 1190 1195 1200 Thr Gly Tyr Arg Leu Asp Pro Gln Lys Ala Gly Ala Leu Val Phe Asn 1205 1210 1215 Ala Leu Trp Glu Glu Gly Ala Val Leu Ser Ala Leu Lys Asn Ala Arg 1220 1225 1230 Phe Ala His Asn Leu Thr Ala Gln Arg Met Glu Phe Asp Tyr Ser Thr 1235 1240 1245 Asn Val Trp Gly Phe Ala Phe Gly Gly Phe Arg Thr Leu Ser Ala Glu 1250 1255 1260 Asn Leu Val Ala Ile Asp Gly Tyr Lys Gly Ala Tyr Gly Gly Ala Ser 1265 1270 1275 Ala Gly Val Asp Ile Gln Leu Met Glu Asp Phe Val Leu Gly Val Ser 1285 1290 1295 Gly Ala Ala Phe Leu Gly Lys Met Asp Ser Gln Lys Phe Asp Ala Glu 1300 1305 1310Val Ser Arg Lys Gly Val Val Gly Ser Val Tyr Thr Gly Phe Leu Ala 1315 1320 1325 Gly Ser Trp Phe Phe Lys Gly Gln Tyr Ser Leu Gly Glu Thr Gln Asn 1330 1335 1340
Asp Met Lys Thr Arg Tyr Gly Val Leu Gly Glu Ser Ser Ala Ser Trp 1345 1350 1355 1360 Thr Ser Arg Gly Val Leu Ala Asp Ala Leu Val Glu Tyr Arg Ser Leu 1365 1370 1375 Val Gly Pro Val Arg Pro Thr Phe Tyr Ala Leu His Phe Asn Pro Tyr 1380 1385 1390 Val Glu Val Ser Tyr Ala Ser Met Lys Phe Pro Gly Phe Thr Glu Gln 1400 1405 Gly Arg Glu Ala Arg Ser Phe Glu Asp Ala Ser Leu Thr Asn Ile Thr 1410 1415 1420 Ile Pro Leu Gly Met Lys Phe Glu Leu Ala Phe Ile Lys Gly Gln Phe 1425 1430 1435 1440 Ser Glu Val Asn Ser Leu Gly Ile Ser Tyr Ala Trp Glu Ala Tyr Arg 1445 1450 1455 Lys Val Glu Gly Gly Ala Val Gln Leu Leu Glu Ala Gly Phe Asp Trp 1460 1465 1470 Glu Gly Ala Pro Met Asp Leu Pro Arg Gln Glu Leu Arg Val Ala Leu 1475 1480 1485 Glu Asn Asn Thr Glu Trp Ser Ser Tyr Phe Ser Thr Val Leu Gly Leu

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385 Ser	Tla	Thr	ሞኮሎ	Pro	390 Pro	T.e.ii	Val	G1 v	Glu	395 Val	Tle	Pha	Ser	Glu	400
				405				_	410					415	
			420		Gly			425					430		
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Pro 465	'Glu	Ser	Ser	Thr	Pro 470	Ser	Ser	Ser	Ser	Pro 475	Ala	Ser	Thr	Pro	Glu 480
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Pro	Ala	Ala	Pro 500	Ser	Leu	Thr	Glu	Ala 505	Glu	Ser	Asp	Gln	Thr 510	Asp	Gln
Thr	Glu	Thr 515	Ser	Asp	Thr	Asn	Ser 520	Asp	Ile	Asp	Val	Ser 525	Ile	Glu	Asn
Ile	Leu 530	Asn	Val	Ala	Ile	Asn 535	Gln	Asn	Thr	Ser	Ala 540	Lys	Lys	Gly	Gly
Ala 545	Ile	Tyr	Gly	Lys	Lys 550	Ala	Lys	Leu	Ser	Arg 555	Ile	Asn	Asn	Leu	Glu 560
Leu	Ser	Gly	Asn	Ser 565	Ser	Gln	Asp	Val	Gly 570	Gly	Gly	Leu	Cys	Leu 575	Thr
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Ile 625	Val	Glu	Ser	Thr	Pro 630	Glu	Ala	Pro	Glu	Glu 635	Ile	Pro	Pro	Val	Glu 640
Gly	Glu	Glu	Ser	Thr 645	Ala	Thr	Glu	Asn	Pro 650	Asn	Ser	Asn	Thr	Glu 655	Gly
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Thr	Gly	Thr 675	Gly	Val	Val	Asn	Asn 680	Glu	Ser	Gln	Asp	Thr 685	Ser	Asp	Thr
Gly	Asn 690	Ala	Glu	Ser	Gly	Glu 695	Gln	Leu	Gln	Asp	Ser 700	Thr	Gln	Ser	Asn
Glu 705	Glu	Asn	Thr	Leu	Pro 710	Asn	Ser	Ser	Ile	Asp 715	Gln	Ser	Asn	Glu	Asn 720
Thr	Asp	Glu	Ser	Ser 725	Asp	Ser	His	Thr	Glu 730	Glu	Ile	Thr	Asp	Glu 735	Ser
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Ala	Cys 770	Leu	Ala	Lys	Ser	Tyr 775	Ala	Ala	Ser	Thr	Asp 780	Ser	Ser	Pro	Val
Ser 785	Asn	Ser	Ser	Gly	Ser 790	Asp	Val	Thr	Ala	Ser 795	Ser	Asp	Asn	Pro	Asp 800
				805	Asp				810					815	
Pro	Glu	Ala	Gly 820	Ser	Thr	Thr	Glu	Thr 825	Pro	Thr	Leu	Ile	Gly 830	Gly	Glÿ
Ala	Ile	Tyr 835	Gly	Glu	Thr	Val	Lys 840		Glu	Asn	Phe	Ser 845		Gln	Gly
Ile	Phe 850		Gly	Asn	Lys	Ala 855	Ile	Asp	Asn	Thr	Thr 860	Glu	Gly	Ser	Ser
Ser 865	Lys	Ser	Asn	Val	Leu 870	Gly	Gly	Ala	Val	Tyr 875		Lys	Thr	Leu	Phe 880

Asn Leu Asp Ser Gly Ser Ser Arg Arg Thr Val Thr Phe Ser Gly Asn 885 890 895 Thr Val Ser Ser Gln Ser Thr Thr Gly Gln Val Ala Gly Gly Ala Ile 905 910. Tyr Ser Pro Thr Val Thr Ile Ala Thr Pro Val Val Phe Ser Lys Asn 920 925 Ser Ala Thr Asn Asn Ala Asn Asn Ala Thr Asp Thr Gln Arg Lys Asp 935 940 Thr Phe Gly Gly Ala Ile Gly Ala Thr Ser Ala Val Ser Leu Ser Gly 950 955 Gly Ala His Phe Leu Glu Asn Val Ala Asp Leu Gly Ser Ala Ile Gly 965 970 975 Leu Val Pro Asp Thr Gln Asn Thr Glu Thr Val Lys Leu Glu Ser Gly 985 990 980 Ser Tyr Tyr Phe Glu Lys Asn Lys Ala Leu Lys Arg Ala Thr Ile Tyr 995 1000 1005 Ala Pro Val Val Ser Ile Lys Ala Tyr Thr Ala Thr Phe Asn Gln Asn 1015 1020 Arg Ser Leu Glu Glu Gly Ser Ala Ile Tyr Phe Thr Lys Glu Ala Ser 1025 1030 1035 Ile Glu Ser Leu Gly Ser Val Leu Phe Thr Gly Asn Leu Val Thr Pro 1045 1050 1055 Thr Leu Ser Thr Thr Thr Glu Gly Thr Pro Ala Thr Thr Ser Gly Asp 1060 1065 1070 Val Thr Lys Tyr Gly Ala Ala Ile Phe Gly Gln Ile Ala Ser Ser Asn 1075 1080 1085 Gly Ser Gln Thr Asp Asn Leu Pro Leu Lys Leu Ile Ala Ser Gly Gly 1090 1095 1100 Asn Ile Cys Phe Arg Asn Asn Glu Tyr Arg Pro Thr Ser Ser Asp Thr 1105 1110 1115 Gly Thr Ser Thr Phe Cys Ser Ile Ala Gly Asp Val Lys Leu Thr Met 1125 1130 1135 Gln Ala Ala Lys Gly Lys Thr Ile Ser Phe Phe Asp Ala Ile Arg Thr 1140 1145 1150 Ser Thr Lys Lys Thr Gly Thr Gln Ala Thr Ala Tyr Asp Thr Leu Asp 1155 1160 1165 Ile Asn Lys Ser Glu Asp Ser Glu Thr Val Asn Ser Ala Phe Thr Gly 1170 1175 1180 Thr Ile Leu Phe Ser Ser Glu Leu His Glu Asn Lys Ser Tyr Ile Pro 1195 1185 1190 Gln Asn Val Val Leu His Ser Gly Ser Leu Val Leu Lys Pro Asn Thr 1205 1210 1215 Glu Leu His Val Ile Ser Phe Glu Gln Lys Glu Gly Ser Ser Leu Val 1220 1225 1230 Met Thr Pro Gly Ser Val Leu Ser Asn Gln Thr Val Ala Asp Gly Ala 1235 1240 1245 Leu Val Ile Asn Asn Met Thr Ile Asp Leu Ser Ser Val Glu Lys Asn 1250 1255 1260 Gly Ile Ala Glu Gly Asn Ile Phe Thr Pro Pro Glu Leu Arg Ile Ile 1270 1275 1280 Asp Thr Thr Thr Ser Gly Ser Gly Gly Thr Pro Ser Thr Asp Ser Glu 1285 1290 1295 Ser Asn Gln Asn Ser Asp Asp Thr Lys Glu Gln Asn Asn Asn Asp Ala 1300 1305 1310 Ser Asn Gln Gly Glu Ser Ala Asn Gly Ser Ser Ser Pro Ala Val Ala 1320 1325 Ala Ala His Thr Ser Arg Thr Arg Asn Phe Ala Ala Ala Ala Thr Ala 1330 1335 1340 Thr Pro Thr Thr Thr Pro Thr Ala Thr Thr Thr Thr Ser Asn Gln Val 1350 1355 Ile Leu Gly Gly Glu Ile Lys Leu Ile Asp Pro Asn Gly Thr Phe Phe

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1585 1590 1595 1600

Thr His Tyr Pro Thr Ile Arg Glu Arg Asn Gln Gly Glu Trp Glu Asp
1605 1610 1615

Leu Gly Trp Leu Thr Ala Leu Arg Val Ser Ser Val Leu Arg Thr Pro 1620 1625 1630 Ala Gln Gly Asp Thr Lys Arg Ile Thr Val Tyr Gly Glu Leu Glu Tyr 1635 1640 1645 Ser Ser Ile Arg Gln Lys Gln Phe Thr Glu Thr Glu Tyr Asp Pro Arg 1650 1655 . 1660 Tyr Phe Asp Asn Cys Thr Tyr Arg Asn Leu Ala Ile Pro Met Gly Leu 1665 1670 1675 Ala Phe Glu Gly Glu Leu Ser Gly Asn Asp Ile Leu Met Tyr Asn Arg 1690 1685 1695 Phe Ser Val Ala Tyr Met Pro Ser Ile Tyr Arg Asn Ser Pro Thr Cys 1700 1705 1710 Lys Tyr Gln Val Leu Ser Ser Gly Glu Gly Glu Ile Ile Cys Gly 1715 1720 1725 Val Pro Thr Arg Asn Ser Ala Arg Gly Glu Tyr Ser Thr Gln Leu Tyr 1730 1735 1740 Pro Gly Pro Leu Trp Thr Leu Tyr Gly Ser Tyr Thr Ile Glu Ala Asp 1745 1750 1755 1760 Ala His Thr Leu Ala His Met Met Asn Cys Gly Ala Arg Met Thr Phe 1765 1770

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<211> 1752

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So			35					40					45			
65 70 75 80 80 85 87 81 80 85 Thr 70 90 90 95 Thr 75 85 77 95 75 85 Thr 85 77 95 75 85 77 90 95 75 85 77 75 85 77 75 85 77 75 85 77 75 85 77 75 85 77 75 85 75 110	_	50					55	_				60		_		•
S	65					70					75					80
Ser				85					90		_			95		
115	Thr	Thr	Thr		Asp	Pro	Lys	Gly		Gly	Ala	Phe	Tyr		Ala	His
130	Ser	Gly		Leu	Ser	Phe	Met		Arg	Ser	Gly	Thr		Gly	Ser	Leu
150	Thr		Ser	Glu	Ile	Lys		Thr	Gly	Glu	Gly		Ala	Ile	Phe	Ser
Ser Leu Ser Gly Tile Thr Lys Ala Thr Phe Ser Cys Asn Ser Ala Glu 190		Gly	Glu	Leu	Leu		Thr	Asp	Leu	Thr		Leu	Thr	Ile	Gln	
180	Asn	Leu	Ser	Gln		Ser	Gly	Gly	Ala		Phe	Gly	Gly	Ser		Ile
Ser Glu Thr Ser Gly Ser Ser Ser Ser Ser Gly Asp Asp Ser Val Ser Zer Ser	Leu	Ser		Ile	Thr	Lys	Ala		Phe	Ser	Суѕ	Asn		Ala	Glu	
Ser	Val	Pro		Pro	Val	Lys	Lys		Thr	Glu	Pro	Lys		Gln	Thr	Ala
225	Ser		Thr	Ser	Gly	Ser		Ser	Ser	Ser	Gly		Asp	Ser	Val	Ser
The Ser The Pro Ser His Lys Pro Gly Ser Gly Gly Ala Ile Tyr Ala 265 Lys Gly Asp Leu Thr Ile Ala Asp Ser Gln Glu Val Leu Phe Ser Ile 275 Asn Lys Ala Thr Lys Asp Gly Gly Ala Ile Phe Ala Glu Lys Asp Val 295 Ser Plo Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu 305 Glu Lys Gly Gly Ala Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu 305 Ser Lys Gln Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser Ile Gln Ser 330 Ser Lys Gln Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser Ile Gln Ser 335 Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln Gly Gly Gly Gly Ala Ile 355 Ala Leu Tyr Val Glu Gly Gly Gly Ile Asn Phe Glu Thr Lys Lys Ile Thr 355 Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 370 Ala Ser Ser Leu Lys Ala Gln Ala Ser Ala Gly Asn Asn Asn Asn Lys Ala Trp 385 Arg Ile Lys Tyr Asn Ser Gly Ser Gly Ser Asp Ser Asn Leu Glu Thr Val 420 Ala Ser Ser Ser Ser Ser Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser 445 Fer Val Thr Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr 440 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Gly Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Gly Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Gly Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Gly Asn 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly		Pro	Ser	Ser	Ser		Ala	Glu	Pro	Ala		Ala	Asn	Leu	Gln	
Lys Gly Asp Leu Thr Ile Ala Asp Ser Gln Glu Val Leu Phe Ser Ile 270 Asn Lys Ala Thr Lys Asp Gly Gly Ala Ile Phe Ala Glu Lys Asp Val 290 Ser Phe Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu 305 Glu Lys Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser Ile 325 Ser Lys Gln Ser Leu Phe Ala Glu Ser	His	Phe	Ile	Cys		Thr	Ala	Thr	Pro		Ala	Gln	Thr	Asp		Glu
Asn Lys Ala Thr Lys Asp Gly Gly Ala Ile Phe Ala Glu Lys Asp Val 295 Ser Phe Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu 305 Ser Phe Gly Gly Ala Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu 305 Ser Lys Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser Ile Gln Ser 325 Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln Gly Gly Gly Gly 345 Ala Leu Tyr Val Glu Gly Gly Ile Asn Phe Gln Asp Leu Glu Glu Glu Ile 355 Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Gln Thr Lys Lys Ile Thr 370 Leu Pro Ser Leu Lys Ala Gln Ala Ser Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 385 Leu Pro Ser Leu Lys Ala Gln Ala Ser Ala Gly Ala Gly Asn Ala Asp Ala Trp 385 Ser Gly Asp Ser Ser Ser Gly Ser Asp Ser Asp Thr Ser Glu Thr Val 415 Ser Gly Asp Ser Ser Ser Gly Ser Asp Ser Asp Thr Ser Glu Thr Val 420 Pro Val Thr Ala Lys Gly	Thr	Ser	Thr		Ser	His	Lys	Pro		Ser	Gly	Gly	Ala		Tyr	Ala
Ser Phe Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu 305 Glu Lys Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser Ile Gln Ser 320 Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln Gly Gly Gly Gly 340 Ala Leu Tyr Val Glu Gly Gly Ile Asn Phe Gln Asp Leu Glu Glu Ile 360 Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 370 Leu Pro Ser Leu Lys Ala Gln Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 380 Ala Ser Ser Ser Pro Gln Ser Gly Ser Gly Ala Thr Thr Val Ser Asp 400 Ala Ser Ser Ser Ser Fro Gln Ser Gly Ser Asp Thr Ser Glu Thr Lys Lys Ala Trp 385 Ser Gly Asp Ser Ser Gly Ser Asp Thr Ser Glu Thr Val Ser Asp 425 Pro Val Thr Ala Lys Gly Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser 435 Ile Thr Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr 465 Asp Val Gly Gly Gly Ala Tyr Val Lys Asn Ser Ser Ser Asp Lys Glu Asn 465 Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Ser Asp Lys Gln Gly Gly Lys Gly Ile Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asp Lys Gln Gly Gly Lys	Lys	Gly		Leu	Thr	Ile	Ala		Ser	Gln	Glu	Val		Phe	Ser	Ile
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Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln Gly Gly Gly Gly 340 Ala Leu Tyr Val Glu Gly Gly Ile Asn Phe Gln Asp Leu Glu Glu Ile 355 Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 370 Leu Pro Ser Leu Lys Ala Gln Ala Ser Ala Gly Asn Ala Asp Ala Trp 385 Ala Ser Ser Ser Pro Gln Ser Gly Ser Gly Ala Thr Thr Val Ser Asp 400 Ala Ser Gly Asp Ser Ser Ser Gly Ser Gly Ser Asp Thr Ser Glu Thr Val Ser Asp 415 Ser Gly Asp Ser Ser Ser Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser Asp 425 Ile Thr Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr 450 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr 465 Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gln Gly Gly Gly Ile Ile Thr Lys Asn Asn Lys Gly Gly Gly Ile Ile Ile Glu Ile Ala Asn Lys Gly Gly Gly Gly Gly Ile Ile Ile Ile Gly Ile Ile Ile Ile Gly Ile		Phe	Glu	Asn	Ile		Ser	Leu	Lys	Val		Thr	Asn	Gly	Ala	
Ala Leu Tyr Val Glu Gly Gly Ile Asn Phe Gln Asp Leu Glu Glu Ile 365 Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 370 Leu Pro Ser Leu Lys Ala Gln Ala Ser Ala Gly Asn Ala Asp Ala Trp 385 Ala Ser Ser Ser Pro Gln Ser Gly Ser Gly Ala Thr Thr Val Ser Asp 400 Ala Ser Ser Ser Ser Ser Gly Ser Gly Ser Asp Thr Ser Glu Thr Val Ser Asp 405 Ser Gly Asp Ser Ser Ser Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser 445 Ile Thr Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr 450 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Gly Thr Leu Thr Cys Glu Asn 465 Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gly Gly Gly Gly Ile Ile Thr Lys Asn Leu Ser 480 Gly Ile Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asp Lys Gly Gly Gly Gly Ile Ile Ile Ser Asp Lys Gly Gly Gly Gly Gly Gly Gly Gly Asn Asn Leu Thr Gly Ile Ile Ile Ser Asp Ser Asp Lys Glu Asn 480 Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gln Gly Gly Gly Ile Ile Tyr Gly Lys	Glu	Lys	Gly	Gly		Ile	Tyr	Ala	Lys		Asp	Leu	Ser	Ile		Ser
Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 370				340					345			_		350	_	-
370	Ala	Leu		Val	Glu	Gly	Gly		Asn	Phe	Gln	Asp		Glu	Glu	Ile
385	_	370					375					380	_	_		
Ser Gly Asp Ser Ser Ser Gly Ser Asp Ser Asp Thr Ser Glu Thr Val 420	385					390					395			_		400
Pro Val Thr Ala Lys Gly Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser 435 Ile Thr Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr 450 Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn 465 Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gln Gly Gly 485 Gly Ile Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asn Leu Thr Gly Lys	Ala	Ser	Ser	Ser		Gln	Ser	Gly	Ser		Ala	Thr	Thr	Val		Asp
Asp Val Gly Gly Gln Phe Leu Lys Asn Ser Asp Lys Gln Gly Gly Gly Asp Ass Ser His Arg Leu Gln Phe Leu Lys Asn Asn Lys Gly Gly Gly Gly Gly Asn Asn Lys Gly Gly Gly Gly Asn Asn Asn Lys Gly Gly Asn Asn Asn Lys Glu Asn Asn Asn Lys Glu Asn Asn Asn Asn Lys Glu Asn Asn Asn Asn Asn Lys Glu Asn	Ser	Gly	Asp		Ser	Ser	Gly	Ser		Ser	Asp	Thr	Ser		Thr	Val
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Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gln Gly Gly 485 490 495 Gly Ile Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asn Leu Thr Gly Lys		Val	Gly	Gly	Gly		Tyr	Val	Lys	Gly		Leu	Thr	Суѕ	Glu	
Gly Ile Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asn Leu Thr Gly Lys		His	Arg	Leu			Leu	Lys	Asn			Asp	Lys	Gln		
	Gly	Ile	Tyr		Glu	Asp	Asn	Ile		Leu	Ser	Asn	Leu			Lys

Thr	Leu	Phe 515	Gln	Glu	Asn	Thr	Ala 520	Lys	Glu	Glu	Gly	Gly 525	Gly	Leu	Phe
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545				Asn	550					555					560
	_			Ser 565					570					575	
			580	Val				585					590		
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625	_			Thr	630		-			635					640
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			660	Ala				665					670	•	
		675		Ala			680					685			
	690		•	Asp		695					700				
705				Thr	710					715					720
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-			900	Gly				905					910		
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945				Asp	950					955		_	•		960
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	530					535	Ser				540				
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-				565			Asn		570	-			•	575	-
			580				Ile	585					590		
		595					Ser 600					605			
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625	_			_	630		Thr Leu			635					640
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			660				Ser	665					670		
		675					680 Tyr					685			
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705					710		Gly			715					720
				725			Gly		730					735	•
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	_	755		_			760 Glu					765			
	770					775	Tyr				780				
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		_		805			Arg		810					815	
			820		•			825	-			,	830		

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Phe	Asp	Gly	Asn 420		Lys	Arg	Thr	Ala 425		Glu	Asn	Ala	Ala 430	Asp	Val
Asn	Gly	Val 435		Val	Ser	Ser	Gln 440		Ile	Ser	Met	Gly 445	Ser	Gly	Gly
Lys	Ile 450		Thr	Leu	Arg	Ala 455		Ala	Gly	His	Gln 460	Ile	Leu	Phe	Asn
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				805					810					815	
_			820		Ala			825					830		
		835			Ala		840					845			
-	850				Ile	855					860				
Ser	Lys	ьeu	Tyr	ьeu	Asn	GTI	ьeu	Arg	PIO	rne	val	GTU	нта	GTU	rue

870

WO 02/08267 PCT/US01/23121

Ser Tyr Ala Asp His Glu Ser Phe Thr Glu Glu Gly Asp Gln Ala Arg 885 890 Ala Phe Lys Ser Gly His Leu Leu Asn Leu Ser Val Pro Val Gly Val 905 Lys Phe Asp Arg Cys Ser Ser Thr His Pro Asn Lys Tyr Ser Phe Met 915 920 925 Ala Ala Tyr Ile Cys Asp Ala Tyr Arg Thr Ile Ser Gly Thr Glu Thr 935 Thr Leu Leu Ser His Gln Glu Thr Trp Thr Thr Asp Ala Phe His Leu 945 950 955 960 Ala Arg His Gly Val Val Val Arg Gly Ser Met Tyr Ala Ser Leu Thr 965 970 975 965 Ser Asn Ile Glu Val Tyr Gly His Gly Arg Tyr Glu Tyr Arg Asp Ala 980 985 990 Ser Arg Gly Tyr Gly Leu Ser Ala Gly Ser Lys Val Arg Phe <210> 191 <211> 977 <212> PRT <213> Chlamydia <400> 191 Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Asp Ser Ser Leu 10 Val Pro Ser Ser Asp Pro His His His His His Gly Leu Ala Arg 20 25 Glu Val Pro Ser Arg Ile Phe Leu Met Pro Asn Ser Val Pro Asp Pro 35 40 45 Thr Lys Glu Ser Leu Ser Asn Lys Ile Ser Leu Thr Gly Asp Thr His 55 60 Asn Leu Thr Asn Cys Tyr Leu Asp Asn Leu Arg Tyr Ile Leu Ala Ile
70 75 80 70 75 Leu Gln Lys Thr Pro Asn Glu Gly Ala Ala Val Thr Ile Thr Asp Tyr 85 90 Leu Ser Phe Phe Asp Thr Gln Lys Glu Gly Ile Tyr Phe Ala Lys Asn 105 Leu Thr Pro Glu Ser Gly Gly Ala Ile Gly Tyr Ala Ser Pro Asn Ser 120 115 125 Pro Thr Val Glu Ile Arg Asp Thr Ile Gly Pro Val Ile Phe Glu Asn 130 135 140 Asn Thr Cys Cys Arg Leu Phe Thr Trp Arg Asn Pro Tyr Ala Ala Asp 150 ⁻ 155 Lys Ile Arg Glu Gly Gly Ala Ile His Ala Gln Asn Leu Tyr Ile Asn 165 170 175 165 170 175 His Asn His Asp Val Val Gly Phe Met Lys Asn Phe Ser Tyr Val Gln 185 190 Gly Gly Ala Ile Ser Thr Ala Asn Thr Phe Val Val Ser Glu Asn Gln 195 200 205 Ser Cys Phe Leu Phe Met Asp Asn Ile Cys Ile Gln Thr Asn Thr Ala 215 220 Gly Lys Gly Gly Ala Ile Tyr Ala Gly Thr Ser Asn Ser Phe Glu Ser 230 235 240 Asn Asn Cys Asp Leu Phe Phe Ile Asn Asn Ala Cys Cys Ala Gly Gly 245 250 Ala Ile Phe Ser Pro Ile Cys Ser Leu Thr Gly Asn Arg Gly Asn Ile 265 Val Phe Tyr Asn Asn Arg Cys Phe Lys Asn Val Glu Thr Ala Ser Ser 280 285 Glu Ala Ser Asp Gly Gly Ala Ile Lys Val Thr Thr Arg Leu Asp Val

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305					310					315					320
Tyr	Gly	Gly	Ala	Ile 325	Tyr	Ala	Pro	Val	Val 330	Thr	Leu	Val	Asp	Asn 335	Gly
Pro	Thr	Tyr	Phe 340	Ile	Asn	Asn	Ile	Ala 345	Asn	Asn	Lys	Gly	Gly 350	Ala	Ile
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Ile	Phe	Tyr	Asp 420		Ile	Glu	Val	Ser 425		Ala	Gly	Val	Ser 430		Ser
Phe	Asn	Lys 435	Glu	Ala	Asp	Gln	Thr 440		Ser	Val	Val	Phe 445		Gly	Ala
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Pro 465	Ala	Pro	Leu	Thr	Leu 470	Ser	Asn	Gly	Phe	Leu 475	Cys	Ile	Glu	Asp	His 480
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			Gly 500					505					510	_	_
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			Asp	645					650			-		655	
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	690		Asn			695					700				
705			Gly		710					715					720
			Pro	725					730					735	_
			Gly 740					745					750		
		755	Leu				760					765			_
Asn	Tyr 770	ser	Суѕ	GIN	G⊥λ	Glu 775	Met	Leu	Phe	ser	Leu 780	Gln	Glu	Gly	Phe

PCT/US01/23121

Leu Leu Thr Lys Leu Val Gly Leu Tyr Ser Tyr Gly Asp His Asn Cys 795 His His Phe Tyr Thr Gln Gly Glu Asn Leu Thr Ser Gln Gly Thr Phe 805 810 Arg Ser Gln Thr Met Gly Gly Ala Val Phe Phe Asp Leu Pro Met Lys 820 825 830 Pro Phe Gly Ser Thr His Ile Leu Thr Ala Pro Phe Leu Gly Ala Leu 835 840 Gly Ile Tyr Ser Ser Leu Ser His Phe Thr Glu Val Gly Ala Tyr Pro 850 855 860 Arg Ser Phe Ser Thr Lys Thr Pro Leu Ile Asn Val Leu Val Pro Ile 865 870 875 880 Gly Val Lys Gly Ser Phe Met Asn Ala Thr His Arg Pro Gln Ala Trp 885 890 Thr Val Glu Leu Ala Tyr Gln Pro Val Leu Tyr Arg Gln Glu Pro Gly 905 Ile Ala Thr Gln Leu Leu Ala Ser Lys Gly Ile Trp Phe Gly Ser Gly 915 920 925 Ser Pro Ser Ser Arg His Ala Met Ser Tyr Lys Ile Ser Gln Gln Thr 940 930 - 935 Gln Pro Leu Ser Trp Leu Thr Leu His Phe Gln Tyr His Gly Phe Tyr 945 950 955 Ser Ser Ser Thr Phe Cys Asn Tyr Leu Asn Gly Glu Ile Ala Leu Arg 970

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Ala Phe Gly Gly Asp Ile Val Phe Lys Gly Asn Ser Ser Phe Arg Ala 230 235 Gln Gly Ser Asp Ala Ile Tyr Phe Ala Gly Lys Glu Ser His Ile Thr 245 250 Ala Leu Asn Ala Thr Glu Gly His Ala Ile Val Phe His Asp Ala Leu 260 265 Val Phe Glu Asn Leu Lys Glu Arg Lys Ser Ala Glu Val Leu Leu Ile 280 285 Asn Ser Arg Glu Asn Pro Gly Tyr Thr Gly Ser Ile Arg Phe Leu Glu 295 300 Ala Glu Ser Lys Val Pro Gln Cys Ile His Val Gln Gln Gly Ser Leu 315 310 Glu Leu Leu Asn Gly Ala Thr Leu Cys Ser Tyr Gly Phe Lys Gln Asp 325 330 Ala Gly Ala Lys Leu Val Leu Ala Ala Gly Ser Lys Leu Lys Ile Leu 340 345 Asp Ser Gly Thr Pro Val Gln Gly His Ala Ile Ser Lys Pro Glu Ala 360 355 365 Glu Ile Glu Ser Ser Ser Glu Pro Glu Gly Ala His Ser Leu Trp Ile 370 375 Ala Lys Asn Ala Gln Thr Thr Val Pro Met Val Asp Ile His Thr Ile 390 395 Ser Val Asp Leu Ala Ser Phe Ser Ser Gln Gln Glu Gly Thr Val 405 410 Glu Ala Pro Gln Val Ile Val Pro Gly Gly Ser Tyr Val Arg Ser Gly 420 425 Glu Leu Asn Leu Glu Leu Val Asn Thr Thr Gly Thr Gly Tyr Glu Asn 435 440 His Ala Leu Leu Lys Asn Glu Ala Lys Val Pro Leu Met Ser Phe Val 455 460 Ala Ser Ser Asp Glu Ala Ser Ala Glu Ile Ser Asn Leu Ser Val Ser 470 475 Asp Leu Gln Ile His Val Ala Thr Pro Glu Ile Glu Glu Asp Thr Tyr 485 490 495 Gly His Met Gly Asp Trp Ser Glu Ala Lys Ile Gln Asp Gly Thr Leu 500 505 Val Ile Asn Trp Asn Pro Thr Gly Tyr Arg Leu Asp Pro Gln Lys Ala 520 525 Gly Ala Leu Val Phe Asn Ala Leu Trp Glu Glu Gly Ala Val Leu Ser 530 535 540 Ala Leu Lys Asn Ala Arg Phe Ala His Asn Leu Thr Ala Gln Arg Met 545 550 555 Glu Phe Asp Tyr Ser Thr Asn Val Trp Gly Phe Ala Phe Gly Gly Phe 565 570 575 Arg Thr Leu Ser Ala Glu Asn Leu Val Ala Ile Asp Gly Tyr Lys Gly 585 Ala Tyr Gly Gly Ala Ser Ala Gly Val Asp Ile Gln Leu Met Glu Asp 595 600 605 Phe Val Leu Gly Val Ser Gly Ala Ala Phe Leu Gly Lys Met Asp Ser 615 620 Gln Lys Phe Asp Ala Glu Val Ser Arg Lys Gly Val Val Gly Ser Val 630 635 Tyr Thr Gly Phe Leu Ala Gly Ser Trp Phe Phe Lys Gly Gln Tyr Ser 650 Leu Gly Glu Thr Gln Asn Asp Met Lys Thr Arg Tyr Gly Val Leu Gly 660 665 Glu Ser Ser Ala Ser Trp Thr Ser Arg Gly Val Leu Ala Asp Ala Leu 680 Val Glu Tyr Arg Ser Leu Val Gly Pro Val Arg Pro Thr Phe Tyr Ala 695 700 Leu His Phe Asn Pro Tyr Val Glu Val Ser Tyr Ala Ser Met Lys Phe

710 Pro Gly Phe Thr Glu Gln Gly Arg Glu Ala Arg Ser Phe Glu Asp Ala 725 730 Ser Leu Thr Asn Ile Thr Ile Pro Leu Gly Met Lys Phe Glu Leu Ala 745 740 750 Phe Ile Lys Gly Gln Phe Ser Glu Val Asn Ser Leu Gly Ile Ser Tyr 760 Ala Trp Glu Ala Tyr Arg Lys Val Glu Gly Gly Ala Val Gln Leu Leu 770 780 Glu Ala Gly Phe Asp Trp Glu Gly Ala Pro Met Asp Leu Pro Arg Gln 785 790 795 800 795 Glu Leu Arg Val Ala Leu Glu Asn Asn Thr Glu Trp Ser Ser Tyr Phe 810 805 Ser Thr Val Leu Gly Leu Thr Ala Phe Cys Gly Gly Phe Thr Ser Thr 825 Asp Ser Lys Leu Gly Tyr Glu Ala Asn Thr Gly Leu Arg Leu Ile Phe <210> 193 <211> 778 <212> PRT <213> Chlamydia <400> 193 Met His His His His His Gly Leu Ala Ser Cys Val Asp Leu His 5 10 Ala Gly Gly Gln Ser Val Asn Glu Leu Val Tyr Val Gly Pro Gln Ala 20 25 Val Leu Leu Asp Gln Ile Arg Asp Leu Phe Val Gly Ser Lys Asp 35 40 45 Ser Gln Ala Glu Gly Gln Tyr Arg Leu Ile Val Gly Asp Pro Ser Ser 50 60Phe Gln Glu Lys Asp Ala Asp Thr Leu Pro Gly Lys Val Glu Gln Ser 65 70 75 80 Thr Leu Phe Ser Val Thr Asn Pro Val Val Phe Gln Gly Val Asp Gln 90 Gln Asp Gln Val Ser Ser Gln Gly Leu Ile Cys Ser Phe Thr Ser Ser 105 100 110 Ash Leu Asp Ser Pro Arg Asp Gly Glu Ser Phe Leu Gly Ile Ala Phe 115 120 125 Val Gly Asp Ser Ser Lys Ala Gly Ile Thr Leu Thr Asp Val Lys Ala 135 Ser Leu Ser Gly Ala Ala Leu Tyr Ser Thr Glu Asp Leu Ile Phe Glu 150 155 Lys Ile Lys Gly Gly Leu Glu Phe Ala Ser Cys Ser Ser Leu Glu Gln 165 170 175 Gly Gly Ala Cys Ala Ala Gln Ser Ile Leu Ile His Asp Cys Gln Gly 185 190 180 Leu Gln Val Lys His Cys Thr Thr Ala Val Asn Ala Glu Gly Ser Ser 195 200 Ala Asn Asp His Leu Gly Phe Gly Gly Gly Ala Phe Phe Val Thr Gly 215 220 Ser Leu Ser Gly Glu Lys Ser Leu Tyr Met Pro Ala Gly Asp Met Val 225 230 235 240 Val Ala Asn Cys Asp Gly Ala Ile Ser Phe Glu Gly Asn Ser Ala Asn
245
255 245 250 Phe Ala Asn Gly Gly Ala Ile Ala Ala Ser Gly Lys Val Leu Phe Val 265 Ala Asn Asp Lys Lys Thr Ser Phe Ile Glu Asn Arg Ala Leu Ser Gly 275 285 280 Gly Ala Ile Ala Ala Ser Ser Asp Ile Ala Phe Gln Asn Cys Ala Glu

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	370			Ser		375					380				
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				Ser 405					410					415	
			420	Ala				425					430		
		435		Ser			440					445			
-	450			Tyr		455					460				
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	530			Lys		535					540				
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				Glu 565					570					575	
			580					585					590		
		595		Gly			600					605			
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_				645 Leu					650					655	
			660					665					670		Gly
		675					680					685			Glu
-	690					695					700				Asp
705				Ser	710					715					720
				725					730					735	Pro
			740	ı				745					750		Ala
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PCT/US01/23121 WO 02/08267 95

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Val	Glu	Lys 435	Asn	Gly	Ile	Ala	Glu 440	Gly	Asn	Ile	Phe	Thr 445	Pro	Pro	Glu
Leu	Arg 450	Ile	Ile	Asp	Thr	Thr 455	Thr	Ser	Gly	Ser	Gly 460	Gly	Thr	Pro	Ser
Thr 465	Asp	Ser	Glu	Ser	Asn 470	Gln	Asn	Ser	Asp	Asp 475	Thr	Lys	Glu	Gln	Asn 480
Asn	Asn	Asp	Ala	Ser 485	Asn	Gln	Gly	Glu	Ser 490	Ala	Asn	Gly	Ser	Ser 495	Ser
Pro	Ala	Val	Ala 500	Ala	Ala	His		Ser 505	Arg	Thr	Arg	Asn	Phe 510	Ala	Ala
Ala	Ala	Thr 515	Ala	Thr	Pro	Thr	Thr 520	Thr	Pro	Thr	Ala	Thr 525	Thr	Thr	Thr
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785			_		790					Leu 795	_				800
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865	-		_		870			_	•	Pro 875			_		880
				885					890	Ser	_			895	
			900					905		Ala	_		910		
Thr	GIn	ьеи	туr	Pro	стλ	rro	тел	Trp	Tnr	Leu	Tyr	етЛ	ser	ıyr	TUL

97

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Leu	Ser	Leu	Ser 420	Asn	Leu	Lys	Thr	Val 425	Thr	Leu	Thr	Lys	Asn 430	Ser	Ala
Lys	Glu	Ser 435	Gly	Gly	Ala	Ile	Phe 440	Thr	Asp	Leu	Ala	Ser 445	Ile	Pro	Thr
Thr	Asp 450	Thr	Pro	Glu	Ser	Ser 455	Thr	Pro	Ser	Ser	Ser 460	Ser	Pro	Ala	Ser
465					470			-		475			Phe		480
Thr	Ala	Glu	Pro	Ala 485	Ala	Pro	Ser	Leu	Thr 490	Glu	Ala	Glu	Ser	Asp 495	Gln
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		515					520					525	Ser		
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		595					600					605	Asp		
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785		_			790					795			Thr		800
			Ala	805	чта	GIY	Ser	T11T	810	OI U	1 4 5 1.	110	7117	815	116
σ±У	атй	атЪ	820	TTG											
	0> 1 1> 5														

<211> 525 <212> PRT <213> Chlamydia

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Asn Ly	ys	Met	Lys	Ser 485	Arg	Lys	Ala	Суѕ	Gly 490	Val	Ala	Val	Gly	Ala 495	Thr	
Leu I	l.e	Asp	Ala 500	Asp	Lys	Trp	Ser	Ile 505	Thr	Gly	Glu	Ala	Arg 510	Leu	Ile	
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                                                                        51
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cagageggee gettagaace ggaetttact tee
                                                                        33
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<212> PRT
<213> Chlamydia
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Ser Thr Asp Leu
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WO 02/08267

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<223> Made in a lab
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Val Ile Val Gly
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Pro Met Pro Arg
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Glu Ile Val Lys
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Val Trp Glu Tyr
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Lys Lys His Asn
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<210> 230
<211> 20
<212> PRT
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1
Pro Asp Ala Asn
            20
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<211> 20
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Asn Cys Gln Asp Gln Lys Asn Lys Arg Asn Ile Leu Pro Asp Ala Asn
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Leu Ala Lys Val
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Lys Asn Lys Arg Asn Ile Leu Pro Asp Ala Asn Leu Ala Lys Val Phe
                                    10
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Gly Ser Ser Asp
<210> 233
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<223> Made in a lab
<400> 233
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Ile Asp Met Phe
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Met Thr Lys Ala
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<212> PRT
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Phe Gly Ser Ser Asp Pro Ile Asp Met Phe Gln Met Thr Lys Ala Leu
Ser Lys His Ile Val Lys
           20
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<212> PRT
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Tyr Pro Val Glu
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<212> PRT
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Ala Val Pro Lys Tyr Ala Thr Val Gly Ser Pro Tyr Pro Val Glu Ile
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Thr Ala Thr Gly
<210> 238
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<212> PRT
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<220>
<223> Made in a lab
<400> 238
Ala Thr Val Gly Ser Pro Tyr Pro Val Glu Ile Thr Ala Thr Gly Lys
Arg Asp Cys Val
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<210> 239
<211> 20
<212> PRT
<213> Artificial Sequence
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Pro Tyr Pro Val Glu Ile Thr Ala Thr Gly Lys Arg Asp Cys Val Asp
1 5
                                10
Val Ile Ile Thr
<210> 240
<211> 21
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Gln. Leu Pro Cys Glu
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Lys Arg Asp Cys Val Asp Val Ile Ile Thr Gln Gln Leu Pro Cys Glu
Ala Glu Phe Val
<210> 242
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 Ser Asp Pro Ala
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 Thr Thr Pro Thr
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 Asp Gly Lys Leu
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 Val Arg Ser Asp Pro Ala Thr Thr Pro Thr Ala Asp Gly Lys Leu Val
 1 5
 Trp Lys Ile Asp
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Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly
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Gly Asp Lys Cys Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly
<210> 254
<211> 20
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<400> 254
Thr Glu Tyr Pro Leu Leu Ala Asp Pro Ser Phe Lys Ile Ser Glu Ala
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Phe Gly Val Leu
            20
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Ala Leu Arg Ala
<210> 257
<211> 20
<212> PRT
<213> Artificial Sequence
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Ala Phe Gly Val Leu Asn Pro Glu Gly Ser Leu Ala Leu Arg Ala Thr
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Phe Leu Ile Asp
           20
<210> 258
<211> 20
<212> PRT
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Asn Pro Glu Gly Ser Leu Ala Leu Arg Ala Thr Phe Leu Ile Asp Lys
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His Gly Val Ile
           20
<210> 259
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<212> PRT
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<223> Made in a lab
<400> 259
Leu Ala Leu Arg Ala Thr Phe Leu Ile Asp Lys His Gly Val Ile Arg
1
                        10
His Ala Val Ile
           20
<210> 260
<211> 20
<212> PRT
<213> Artificial Sequence
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<220>
<223> Made in a lab
<400> 260
Thr Phe Leu Ile Asp Lys His Gly Val Ile Arg His Ala Val Ile Asn
                                    10
Asp Leu Pro Leu
<210> 261
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 261
Lys His Gly Val Ile Arg His Ala Val Ile Asn Asp Leu Pro Leu Gly
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                    . 10
Arg Ser Ile Asp
            20
<210> 262
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
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1
Glu Leu Arg Ile
            20
<210> 263
<211> 897
<212> DNA
<213> Chlamydia
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<222> 604
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acacagccca acaataaaat ggcaagggta gtaaataaga cgaagggagt ggataagact
                                                                       120
attaaggttg ccaagtctgc tgccgaattg accgcaaata ttttggaaca agctggaggc
                                                                       180
gegggetett cegeacacat tacagettee caagtgteea aaggattagg ggatgegaga
                                                                       240
actifttgtcg ctttagggaa tgcctttaac ggagcgttgc caggaacagt tcaaagtgcg
                                                                       300
caaaqcttct tctctcacat gaaagctgct agtcagaaaa cgcaagaagg ggatgagggg
                                                                       360
ctcacagcag atctttgtgt gtctcataag cgcagagcgg ctgcggctgt ctgtagcatc
                                                                       420
atcggaggaa ttacctacct cgcgacattc ggagctatcc gtccgattct gtttgtcaac
                                                                       480
aaaatgctgg caaaaccgtt tctttcttcc caaactaaag caaatatggg atcttctgtt
                                                                       540
agctatatta tggcggctaa ccatgcagcg tctgtggtgg gtgctggact cgctatcagt
                                                                       600
gcgnaaagag cagattgcga agcccgctgc gctcgtattg cgagagaaga gtcgttactc
                                                                       660
qaaqtqccgg gagaggaaaa tgcttgcgag aagaaagtcg ctggagagaa agccaagacg
                                                                       720
ttcacgcgca tcaagtatgc actcctcact atgctcgaga agtttttgga atgcgttgcc
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840

897

WO 02/08267 PCT/US01/23121

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gacgttttca aattggtgcc gctgcctatt acaatgggta ttcgtgcgat tgtggctgct
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Lys Thr Lys Gly Val Asp Lys Thr Ile Lys Val Ala Lys Ser Ala Ala
       35
                          40
                                            45
Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser
                       55
Ala His Ile Thr Ala Ser Gln Val Ser Lys Gly Leu Gly Asp Ala Arg
                                       75
Thr Val Val Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr
               85
                                  90
                                                      95
Val Gln Ser Ala Gln Ser Phe Phe Ser His Met Lys Ala Ala Ser Gln
          100
                               105
                                                  110
Lys Thr Gln Glu Gly Asp Glu Gly Leu Thr Ala Asp Leu Cys Val Ser
                          120
                                              125
His Lys Arg Arg Ala Ala Ala Val Cys Ser Ile Ile Gly Gly Ile
                      135
                                         140
Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn
                 150
                                      155
Lys Met Leu Ala Lys Pro Phe Leu Ser Ser Gln Thr Lys Ala Asn Met
               165
                                  170
Gly Ser Ser Val Ser Tyr Ile Met Ala Ala Asn His Ala Ala Ser Val
           180
                               185
Val Gly Ala Gly Leu Ala Ile Ser Ala Xaa Arg Ala Asp Cys Glu Ala
                          200
                                            205
Arg Cys Ala Arg Ile Ala Arg Glu Glu Ser Leu Leu Glu Val Pro Gly
                      215
                                         220
Glu Glu Asn Ala Cys Glu Lys Lys Val Ala Gly Glu Lys Ala Lys Thr
                   230
                                       235
Phe Thr Arg Ile Lys Tyr Ala Leu Leu Thr Met Leu Glu Lys Phe Leu
               245
                                 250
Glu Cys Val Ala Asp Val Phe Lys Leu Val Pro Leu Pro Ile Thr Met
                              265
                                                 270
Gly Ile Arg Ala Ile Val Ala Ala Gly Cys Thr Phe Thr Ser Ala Ile
                        280
     275
                                           285
Ile Gly Leu Cys Thr Phe Cys Ala Arg Ala
   290
                       295
<210> 265
<211> 897
<212> DNA
<213> Chlamydia
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<221> misc_feature
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PCT/US01/23121 WO 02/08267 114

60

120

180

240

300

360

420

480

540

600

660

720

780

840

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attaaggttg ccaagtctgc tgccgaattg accgcaaata ttttggaaca agctggaggc
gegggetett eegeacacat tacagettee caagtgteea aaggattagg ggatgegaga
actgttgtcg ctttagggaa tgcctttaac ggagcgttgc caggaacagt tcaaagtgcg
caaagcttct tctctcacat gaaagctgct agtcagaaaa cgcaagaagg ggatgagggg
ctcacagcag atctttgtgt gtctcataag cgcagagcgg ctgcggctgt ctgtagcatc
atcggaggaa ttacctacct cgcgacattc ggagctatcc gtccgattct gtttgtcaac
aaaatgctgg caaaaccgtt tctttcttcc caaactaaag caaatatggg atcttctgtt
agctatatta tggcggctaa ccatgcagcg tctgtggtgg gtgctggact cgctatcagt
gegnaaagag cagattgega agceegetge getegtattg egagagaaga gtegttacte
gaagtgccgg gagaggaaaa tgcttgcgag aagaaagtcg ctggagagaa agccaagacg
ttcacgcgca tcaagtatgc actcctcact atgctcgaga agtttttgga atgcgttgcc
gacgttttca aattggtgcc gctgcctatt acaatgggta ttcgtgcgat tgtggctgct
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<212> PRT
<213> Chlamydia
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Lys Ala Phe Phe Thr Gln Pro Asn Asn Lys Met Ala Arg Val Val Asn
                                25
Lys Thr Lys Gly Met Asp Lys Thr Ile Lys Val Ala Lys Ser Ala Ala
                            40
Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser
                        55
Ala His Ile Thr Ala Ser Gln Val Ser Lys Gly Leu Gly Asp Ala Arg
                    70
                                      75
Thr Val Val Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr
               85
                                   90
Val Gln Ser Ala Gln Ser Phe Phe Ser His Met Lys Ala Ala Ser Gln
            100
                                105
Lys Thr Gln Glu Gly Asp Glu Gly Leu Thr Ala Asp Leu Cys Val Ser
                           120
                                                125
His Lys Arg Arg Ala Ala Ala Ala Val Cys Ser Ile Ile Gly Gly Ile
                        135
   130
                                            140
Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn
                    150
                                        155
Lys Met Leu Ala Lys Pro Phe Leu Ser Ser Gln Thr Lys Ala Asn Met
               165
                                   170
Gly Ser Ser Val Ser Tyr Ile Met Ala Ala Asn His Ala Ala Ser Val
                                185
                                                    190
Val Gly Ala Gly Leu Ala Ile Ser Ala Xaa Arg Ala Asp Cys Glu Ala
                            200
                                                205
Arg Cys Ala Arg Ile Ala Arg Glu Glu Ser Leu Leu Glu Val Pro Gly
                       215
                                            220
Glu Glu Asn Ala Cys Glu Lys Lys Val Ala Gly Glu Lys Ala Lys Thr
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118

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Ala Val Pro Ser Ile Val Asn Ser Thr Gln Arg Cys Tyr Gln Tyr Thr
                           120
Arg Gln Ala Phe Glu Leu Gly Ser Lys Thr Lys Glu Arg Lys Thr Pro
 · 130
                       135
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Gly Glu Tyr Ser Lys Met Leu Leu Thr Arg Gly Asp Tyr Leu Leu Ala
145 150 155 160
Ala Ser Arg Glu Ala Cys Thr Ala Val Gly Ala Thr Thr Tyr Ser Ala
165 170
                                   170
Thr Phe Gly Val Leu Arg Pro Leu Met Leu Ile Asn Lys Leu Thr Ala
                               185
                                                   190
Lys Pro Phe Leu Asp Lys Ala Thr Val Gly Asn Phe Gly Thr Ala Val
                         200
                                              205
Ala Gly Ile Met Thr Ile Asn His Met Ala Gly Val Ala Gly Ala Val
                       215
                                           220
Gly Gly Ile Ala Leu Glu Gln Lys Leu Phe Lys Arg Ala Lys Glu Ser
                   230
                                       235
Leu Tyr Asn Glu Arg Cys Ala Leu Glu Asn Gln Gln Ser Gln Leu Ser
              245
                                 250
Gly Asp Val Ile Leu Ser Ala Glu Arg Ala Leu Arg Lys Glu His Val 260 265 270
Ala Thr Leu Lys Arg Asn Val Leu Thr Leu Leu Glu Lys Ala Leu Glu
        275
                           280
                                               285
Leu Val Val Asp Gly Val Lys Leu Ile Pro Leu Pro Ile Thr Val Ala
                       295
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Cys Ser Ala Ala Ile Ser Gly Ala Leu Thr Ala Ala Ser Ala Gly Ile
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                                     315
Gly Leu Tyr Ser Ile Trp Gln Lys Thr Lys Ser Gly Lys
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<400> 293
tgcaatc
<210> 294
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<213> Chlamydia
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Thr Met Gly Ser Leu Val Gly Arg Gln Ala Pro Asp Phe Ser Gly Lys
Ala Val Val Cys Gly Glu Glu Lys Glu Ile Ser Leu Ala Asp Phe Arg
Gly Lys Tyr Val Val Leu Phe Phe Tyr Pro Lys Asp Phe Thr Tyr Val
Cys Pro Thr Glu Leu His Ala Phe Gln Asp Arg Leu Val Asp Phe Glu
Glu His Gly Ala Val Val Leu Gly Cys Ser Val Asp Asp Ile Glu Thr
His Ser Arg Trp Leu Thr Val Ala Arg Asp Ala Gly Gly Ile Glu Gly
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Thr Glu Tyr Pro Leu Leu Ala Asp Pro Ser Phe Lys Ile Ser Glu Ala Phe Gly Val Leu Asn Pro Glu Gly Ser Leu Ala Leu Arg Ala Thr Phe Leu Ile Asp Lys His Gly Val Ile Arg His Ala Val Ile Asn Asp Leu Pro Leu Gly Arg Ser Ile Asp Glu Glu Leu Arg Ile Leu Asp Ser Leu Ile Phe Phe Glu Asn His Gly Met Val Cys Pro Ala Asn Trp Arg Ser Gly Glu Arg Gly Met Val Pro Ser Glu Glu Gly Leu Lys Glu Tyr Phe Gln Thr Met Asp 195 <210> 295 <211> 181 <212> PRT <213> Chlamydia <400> 295 Lys Gly Gly Lys Met Ser Thr Thr Ile Ser Gly Asp Ala Ser Ser Leu Pro Leu Pro Thr Ala Ser Cys Val Glu Thr Lys Ser Thr Ser Ser Ser Thr Lys Gly Asn Thr Cys Ser Lys Ile Leu Asp Ile Ala Leu Ala Ile Val Gly Ala Leu Val Val Val Ala Gly Val Leu Ala Leu Val Leu Cys 50 60 Ala Ser Asn Val Ile Phe Thr Val Ile Gly Ile Pro Ala Leu Ile Ile Gly Ser Ala Cys Val Gly Ala Gly Ile Ser Arg Leu Met Tyr Arg Ser Ser Tyr Ala Ser Leu Glu Ala Lys Asn Val Leu Ala Glu Gln Arg Leu Arg Asn Leu Ser Glu Glu Lys Asp Ala Leu Ala Ser Val Ser Phe Ile Asn Lys Met Phe Leu Arg Gly Leu Thr Asp Asp Leu Gln Ala Leu Glu Ala Lys Val Met Glu Phe Glu Ile Asp Cys Leu Asp Arg Leu Glu Lys Asn Glu Gln Ala Leu Leu Ser Asp Val Arg Leu Val Leu Ser Ser Tyr

124

Thr Arg Trp Leu Asp 180

<210> 296

<211> 124

<212> PRT

<213> Chlamydia

<400> 296

Ile Tyr Glu Val Met Asn Met Asp Leu Glu Thr Arg Arg Ser Phe Ala $5 \hspace{1.5cm} 10 \hspace{1.5cm} 15$

Val Gln Gln Gly His Tyr Gln Asp Pro Arg Ala Ser Asp Tyr Asp Leu 20 25 30

Pro Arg Ala Ser Asp Tyr Asp Leu Pro Arg Ser Pro Tyr Pro Thr Pro 35 40 45

Pro Leu Pro Ser Arg Tyr Gln Leu Gln Asn Met Asp Val Glu Ala Gly 50 60

Phe Arg Glu Ala Val Tyr Ala Ser Phe Val Ala Gly Met Tyr Asn Tyr 65 70 75 80

Val Val Thr Gln Pro Gln Glu Arg Ile Pro Asn Ser Gln Gln Val Glu 85 90 95

Gly Ile Leu Arg Asp Met Leu Thr Asn Gly Ser Gln Thr Phe Ser Asn 100 105 110

Leu Met Gln Arg Trp Asp Arg Glu Val Asp Arg Glu 115 120

<210> 297

<211> 488

<212> PRT

<213> Chlamydia

<400> 297

Lys Gly Ser Leu Pro Ile Leu Gly Pro Phe Leu Asn Gly Lys Met Gly
10

Phe Trp Arg Thr Ser Ile Met Lys Met Asn Arg Ile Trp Leu Leu Leu

Leu Thr Phe Ser Ser Ala Ile His Ser Pro Val Arg Gly Glu Ser Leu 35 40 45

Val Cys Lys Asn Ala Leu Gln Asp Leu Ser Phe Leu Glu His Leu Leu
50 60

Gln Val Lys Tyr Ala Pro Lys Thr Trp Lys Glu Gln Tyr Leu Gly Trp 65 70 75 80

Asp Leu Val Gln Ser Ser Val Ser Ala Gln Gln Lys Leu Arg Thr Gln 85 90 95

Glu Asn Pro Ser Thr Ser Phe Cys Gln Gln Val Leu Ala Asp Phe Ile 100 105 110

Gly Gly Leu Asn Asp Phe His Ala Gly Val Thr Phe Phe Ala Ile Glu Ser Ala Tyr Leu Pro Tyr Thr Val Gln Lys Ser Ser Asp Gly Arg Phe Tyr Phe Val Asp Ile Met Thr Phe Ser Ser Glu Ile Arg Val Gly Asp 150 Glu Leu Leu Glu Val Asp Gly Ala Pro Val Gln Asp Val Leu Ala Thr Leu Tyr Gly Ser Asn His Lys Gly Thr Ala Ala Glu Glu Ser Ala Ala 185 Leu Arg Thr Leu Phe Ser Arg Met Ala Ser Leu Gly His Lys Val Pro Ser Gly Arg Thr Thr Leu Lys Ile Arg Arg Pro Phe Gly Thr Thr Arg Glu Val Arg Val Lys Trp Arg Tyr Val Pro Glu Gly Val Gly Asp Leu Ala Thr Ile Ala Pro Ser Ile Arg Ala Pro Gln Leu Gln Lys Ser Met Arg Ser Phe Phe Pro Lys Lys Asp Asp Ala Phe His Arg Ser Ser Ser Leu Phe Tyr Ser Pro Met Val Pro His Phe Trp Ala Glu Leu Arg Asn His Tyr Ala Thr Ser Gly Leu Lys Ser Gly Tyr Asn Ile Gly Ser Thr 295 Asp Gly Phe Leu Pro Val Ile Gly Pro Val Ile Trp Glu Ser Glu Gly Leu Phe Arg Ala Tyr Ile Ser Ser Val Thr Asp Gly Asp Gly Lys Ser 330 His Lys Val Gly Phe Leu Arg Ile Pro Thr Tyr Ser Trp Gln Asp Met Glu Asp Phe Asp Pro Ser Gly Pro Pro Pro Trp Glu Glu Phe Ala Lys 360 Ile Ile Gln Val Phe Ser Ser Asn Thr Glu Ala Leu Ile Ile Asp Gln Thr Asn Asn Pro Gly Gly Ser Val Leu Tyr Leu Tyr Ala Leu Leu Ser Met Leu Thr Asp Arg Pro Leu Glu Leu Pro Lys His Arg Met Ile Leu Thr Gln Asp Glu Val Val Asp Ala Leu Asp Trp Leu Thr Leu Leu Glu 425

126

Asn Val Asp Thr Asn Val Glu Ser Arg Leu Ala Leu Gly Asp Asn Met
435 440 445

Glu Gly Tyr Thr Val Asp Leu Gln Val Ala Glu Tyr Leu Lys Ser Phe 450 455 460

Gly Arg Gln Val Leu Asn Cys Trp Ser Lys Gly Asp Ile Glu Leu Ser 465 470 475 480

Thr Pro Ile Pro Leu Phe Gly Phe
485

<210> 298

<211> 140

<212> PRT

<213> Chlamydia

<400> 298

Arg Ile Asp Ile Ser Ser Val Thr Phe Phe Ile Gly Ile Leu Leu Ala

Val Asn Ala Leu Thr Tyr Ser His Val Leu Arg Asp Leu Ser Val Ser 20 25 30

Met Asp Ala Leu Phe Ser Arg Asn Thr Leu Ala Val Leu Leu Gly Leu 35 40 45

Val Ser Ser Val Leu Asp Asn Val Pro Leu Val Ala Ala Thr Ile Gly 50 55 60

Met Tyr Asp Leu Pro Met Asn Asp Pro Leu Trp Lys Leu Ile Ala Tyr 65 70 75 80

Thr Ala Gly Thr Gly Gly Ser Ile Leu Ile Ile Gly Ser Ala Ala Gly 85 90 95

Val Ala Tyr Met Gly Met Glu Lys Val Ser Phe Gly Trp Tyr Val Lys
100 105 110

His Ala Ser Trp Ile Ala Leu Ala Ser Tyr Phe Gly Gly Leu Ala Val 115 120 125

Tyr Phe Leu Met Glu Asn Cys Val Asn Leu Phe Val 130 135 140

<210> 299

<211> 361

<212> PRT

<213> Chlamydia

<400> 299

His Gln Glu Ile Ala Asp Ser Pro Leu Val Lys Lys Ala Glu Gln 15

Ile Asn Gln Ala Gln Gln Asp Ile Gln Thr Ile Thr Pro Ser Gly Leu 20 25 30

Asp Ile Pro Ile Val Gly Pro Ser Gly Ser Ala Ala Ser Ala Gly Ser 35

Ala Ala Gly Ala Leu Lys Ser Ser Asn Asn Ser Gly Arg Ile Ser Leu Leu Leu Asp Asp Val Asp Asn Glu Met Ala Ala Ile Ala Met Gln Gly Phe Arg Ser Met Ile Glu Gln Phe Asn Val Asn Asn Pro Ala Thr Ala Lys Glu Leu Gln Ala Met Glu Ala Gln Leu Thr Ala Met Ser Asp Gln Leu Val Gly Ala Asp Gly Glu Leu Pro Ala Glu Ile Gln Ala Ile Lys 120 Asp Ala Leu Ala Gln Ala Leu Lys Gln Pro Ser Ala Asp Gly Leu Ala Thr Ala Met Gly Gln Val Ala Phe Ala Ala Ala Lys Val Gly Gly 155 Ser Ala Gly Thr Ala Gly Thr Val Gln Met Asn Val Lys Gln Leu Tyr Lys Thr Ala Phe Ser Ser Thr Ser Ser Ser Ser Tyr Ala Ala Ala Leu Ser Asp Gly Tyr Ser Ala Tyr Lys Thr Leu Asn Ser Leu Tyr Ser Glu Ser Arg Ser Gly Val Gln Ser Ala Ile Ser Gln Thr Ala Asn Pro Ala Leu Ser Arg Ser Val Ser Arg Ser Gly Ile Glu Ser Gln Gly Arg Ser Ala Asp Ala Ser Gln Arg Ala Ala Glu Thr Ile Val Arg Asp Ser Gln Thr Leu Gly Asp Val Tyr Ser Arg Leu Gln Val Leu Asp Ser Leu Met 265 Ser Thr Ile Val Ser Asn Pro Gln Ala Asn Gln Glu Glu Ile Met Gln Lys Leu Thr Ala Ser Ile Ser Lys Ala Pro Gln Phe Gly Tyr Pro Ala Val Gln Asn Ser Val Asp Ser Leu Gln Lys Phe Ala Ala Gln Leu Glu 315 Arg Glu Phe Val Asp Gly Glu Arg Ser Leu Ala Glu Ser Gln Glu Asn Ala Phe Arg Lys Gln Pro Ala Phe Ile Gln Gln Val Leu Val Asn Ile Ala Ser Leu Phe Ser Gly Tyr Leu Ser 355

<210> 300 <211> 207 <212> PRT <213> Chlamydia <400> 300 Ser Ser Lys Ile Val Ser Leu Cys Glu Gly Ala Val Ala Asp Ala Arg Met Cys Lys Ala Glu Leu Ile Lys Lys Glu Ala Asp Ala Tyr Leu Phe Cys Glu Lys Ser Gly Ile Tyr Leu Thr Lys Lys Glu Gly Ile Leu Ile Pro Ser Ala Gly Ile Asp Glu Ser Asn Thr Asp Gln Pro Phe Val Leu Tyr Pro Lys Asp Ile Leu Gly Ser Cys Asn Arg Ile Gly Glu Trp Leu Arg Asn Tyr Phe Arg Val Lys Glu Leu Gly Val Ile Ile Thr Asp Ser His Thr Thr Pro Met Arg Arg Gly Val Leu Gly Ile Gly Leu Cys Trp Tyr Gly Phe Ser Pro Leu His Asn Tyr Ile Gly Ser Leu Asp Cys Phe Gly Arg Pro Leu Gln Met Thr Gln Ser Asn Leu Val Asp Ala Leu Ala Val Ala Ala Val Val Cys Met Gly Glu Gly Asn Glu Gln Thr Pro Leu Ala Val Ile Glu Gln Ala Pro Asn Met Val Tyr His Ser Tyr Pro Thr Ser Arg Glu Glu Tyr Cys Ser Leu Arg Ile Asp Glu Thr Glu Asp Leu Tyr Gly Pro Phe Leu Gln Ala Val Thr Trp Ser Gln Glu Lys Lys 200 -<210> 301 <211> 183 <212> PRT <213> Chlamydia <400> 301 Ile Pro Pro Ala Pro Arg Gly His Pro Gln Ile Glu Val Thr Phe Asp Ile Asp Ala Asn Gly Ile Leu His Val Ser Ala Lys Asp Ala Ala Ser

Gly Arg Glu Gln Lys Ile Arg Ile Glu Ala Ser Ser Gly Leu Lys Glu

129

Asp Glu Ile Gln Gln Met Ile Arg Asp Ala Glu Leu His Lys Glu Glu

Asp Lys Gln Arg Lys Glu Ala Ser Asp Val Lys Asn Glu Ala Asp Gly 65 70 75 80

Met Ile Phe Arg Ala Glu Lys Ala Val Lys Asp Tyr His Asp Lys Ile

Pro Ala Glu Leu Val Lys Glu Ile Glu Glu His Ile Glu Lys Val Arg

Gln Ala Ile Lys Glu Asp Ala Ser Thr Thr Ala Ile Lys Ala Ala Ser

Asp Glu Leu Ser Thr Arg Met Gln Lys Ile Gly Glu Ala Met Gln Ala 135

Gln Ser Ala Ser Ala Ala Ala Ser Ser Ala Ala Asn Ala Gln Gly Gly

Pro Asn Ile Asn Ser Glu Asp Leu Lys Lys His Ser Phe Ser Thr Arg 170

Pro Pro Ala Gly Gly Ser Ala 180

<210> 302

<211> 232

<212> PRT

<213> Chlamydia

<400> 302

Met Thr Lys His Gly Lys Arg Ile Arg Gly Ile Gln Glu Thr Tyr Asp

Leu Ala Lys Ser Tyr Ser Leu Gly Glu Ala Ile Asp Ile Leu Lys Gln

Cys Pro Thr Val Arg Phe Asp Gln Thr Val Asp Val Ser Val Lys Leu

Gly Ile Asp Pro Arg Lys Ser Asp Gln Gln Ile Arg Gly Ser Val Ser

Leu Pro His Gly Thr Gly Lys Val Leu Arg Ile Leu Val Phe Ala Ala

Gly Asp Lys Ala Ala Glu Ala Ile Glu Ala Gly Ala Asp Phe Val Gly

Ser Asp Asp Leu Val Glu Lys Ile Lys Gly Gly Trp Val Asp Phe Asp

Val Ala Val Ala Thr Pro Asp Met Met Arg Glu Val Gly Lys Leu Gly 120

Lys Val Leu Gly Pro Arg Asn Leu Met Pro Thr Pro Lys Ala Gly Thr

130

Ile Glu Phe Lys Ala Asp Arg Ala Gly Val Cys Asn Val Gly Val Ala Lys Leu Ser Phe Asp Ser Ala Gln Ile Lys Glu Asn Val Glu Ala Leu 185 Cys Ala Ala Leu Val Lys Ala Lys Pro Ala Thr Ala Lys Gly Gln Tyr Leu Val Asn Phe Thr Ile Ser Ser Thr Met Gly Pro Gly Val Thr Val Asp Thr Arg Glu Leu Ile Ala Leu <210> 303 <211> 238 <212> PRT <213> chlamydia Ile Asn Ser Lys Leu Glu Thr Lys Asn Leu Ile Tyr Leu Lys Leu Lys Ile Lys Lys Ser Phe Lys Met Gly Asn Ser Gly Phe Tyr Leu Tyr Asn Thr Gln Asn Cys Val Phe Ala Asp Asn Ile Lys Val Gly Gln Met Thr Glu Pro Leu Lys Asp Gln Gln Ile Ile Leu Gly Thr Thr Ser Thr Pro Val Ala Ala Lys Met Thr Ala Ser Asp Gly Ile Ser Leu Thr Val Ser Asn Asn Pro Ser Thr Asn Ala Ser Ile Thr Ile Gly Leu Asp Ala Glu Lys Ala Tyr Gln Leu Ile Leu Glu Lys Leu Gly Asp Gln Ile Leu Gly

Val Thr Thr Asp Val Val Lys Thr Ile Ala Glu Leu Arg Lys Gly Lys

Asn Ile Glu Thr Leu Leu Gly Gly Thr Glu Ile Gly Lys Phe Thr Val 165 170 175

Gly Ile Ala Asp Thr Ile Val Asp Ser Thr Val Gln Asp Ile Leu Asp

Lys Ile Thr Thr Asp Pro Ser Leu Gly Leu Leu Lys Ala Phe Asn Asn

Phe Pro Ile Thr Asn Lys Ile Gln Cys Asn Gly Leu Phe Thr Pro Arg

Thr Pro Lys Ser Ser Gly Ser Met Phe Leu Val Ser Ala Asp Ile Ile
180 185 190

131

Ala Ser Arg Met Glu Gly Gly Val Val Leu Ala Leu Val Arg Glu Gly 195 200 205

Asp Ser Lys Pro Tyr Ala Ile Ser Tyr Gly Tyr Ser Ser Gly Val Pro 210 215

Asn Leu Cys Ser Leu Arg Thr Arg Ile Ile Asn Thr Gly Leu 225 230 235

<210> 304

<211> 133

<212> PRT

<213> Chlamydia

<400> 304

Gly Ser Gly Thr Gly Asn Ala Leu Lys Ala Phe Phe Thr Gln Pro Ser

Asn Lys Met Ala Arg Val Val Asn Lys Thr Lys Gly Met Asp Lys Thr 35 40 45

Val Lys Val Ala Lys Ser Ala Ala Glu Leu Thr Ala Asn Ile Leu Glu 50 55 60

Gln Ala Gly Gly Ala Gly Ser Ser Ala His Ile Thr Ala Ser Gln Val 65 70 75 80

Ser Lys Gly Leu Gly Asp Thr Arg Thr Val Val Ala Leu Gly Asn Ala 85 90 95

Phe Asn Gly Ala Leu Pro Gly Thr Val Gln Ser Ala Gln Ser Phe Phe 100 105 110

Ser His Met Lys Ala Ala Ser Gln Lys Thr Gln Glu Gly Asp Glu Gly 115 120 125

Leu Thr Ala Asp Leu 130

<210> 305

<211> 125

<212> PRT

<213> Chlamydia

<400> 305

Met Ala Ser Ile Cys Gly Arg Leu Gly Ser Gly Thr Gly Asn Ala Leu
5 10 15

Lys Ala Phe Phe Thr Gln Pro Ser Asn Lys Met Ala Arg Val Val Asn 20 25 30

Lys Thr Lys Gly Met Asp Lys Thr Val Lys Val Ala Lys Ser Ala Ala 35 40 45

Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser 50 60

Ala His Ile Thr Ala Ser Gln Val Ser Lys Gly Leu Gly Asp Thr Arg Thr Val Val Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr Val Gln Ser Ala Gln Ser Phe Phe Ser His Met Lys Ala Ala Ser Gln Lys Thr Gln Glu Gly Asp Glu Gly Leu Thr Ala Asp Leu 120 <210> 306 <211> 38 <212> DNA <213> Chlamydia trachomatis <400> 306 gagageggee geteatgttt ataacaaagg aacttatg 38 <210> 307 <211> 39 <212> DNA <213> Chlamydia trachomatis <400> 307 gagageggee gettaettag gtgagaagaa gggagttte 39 <210> 308 <211> 1860 <212> DNA <213> Chlamydia trachomatis <400> 308 atgeateace ateaceatea caeggeegeg teegataact teeagetgte ceagggtggg 60 cagggattcg ccattccgat cgggcaggcg atggcgatcg cgggccagat caagcttccc 120 accettcata tegggectae egectteete ggettgggtg ttgtegacaa caacggcaac 180 ggcgcacgag tccaacgcgt ggtcgggagc gctccggcgg caagtctcgg catctccacc 240 ggcgacgtga tcaccgcggt cgacggcgct ccgatcaact cggccaccgc gatggcggac 300 gegettaacg ggcatcatec eggtgacgtc atcteggtga cetggcaaac caagteggge 360 ggcacgcgta cagggaacgt gacattggcc gagggacccc cggccgaatt ctgcagatat ccatcacact ggcggccgct catgtttata acaaaggaac ttatgaatcg agttatagaa 420 480 atccatgete actacgatea aagacaactt teteaatete caaatacaaa ettettagta 540 catcatcett atettactet tatteceaag tttetactag gagetetaat egtetatget ccttattcgt ttgcagaaat ggaattagct atttctggac ataaacaagg taaagatcga 660 gatacettta ccatgatete tteetgteet gaaggeacta attacateat caategeaaa 720 ctcatactca gtgatttctc gttactaaat aaagittcat cagggggagc ctttcggaat 780 ctagcaggga aaatttcctt cttaggaaaa aattcttctg cgtccattca ttttaaacac 840 attaatatca atggttttgg agccggagtc ttttctgaat cctctattga atttactgat 900 ttacgaaaac ttgttgcttt tggatctgaa agcacaggag gaatttttac tgcgaaagag 960 gacatetett ttaaaaacaa ecaecacatt geetteegea ataatateae caaagggaat 1020 ggtggcgtta tccagctcca aggagatatg aaaggaagcg tatcctttgt agatcaacgt ggagctatca tctttaccaa taaccaagct gtaacttctt catcaatgaa acatagtggt 1080 1140 cgtggaggag caattagcgg tgacttcgca ggatccagaa ttcttttct taataaccaa 1200 caaattactt tcgaaggcaa tagcgctgtg catggaggtg ctatctacaa taagaatggc 1260 cttgtcgagt tcttaggaaa tgcaggacct cttgccttta aagagaacac aacaatagct 1320 aacgggggag ctatatacac aagtaatttc aaagcgaatc aacaaacatc ccccattcta 1380 ttctctcaaa atcatgcgaa taagaaaggc ggagcgattt acgcgcaata tgtgaactta 1440

atcacctett etcaatgete aattactget cataatacea teaetittie egataatget geeggagate tiggaggagg ageaattett etagaaggga aaaaacette tetaacetig attgeteata giggtaatat tigeatitage gigeaatacea tigeteestata eaceaaaaaa getteeetag ategacacaa tietatetta ateaaagaag eteeetataa aateeaaeti geagegaaca aaaaceatte tatteatte titgateetig teatggeatt gieageatea 18														1500 1560 1620 1680 1740 1800 1860		
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)> 30 His		His	His 5	His	His	Thx	Ala	Ala 10	Ser	Asp	Asn	Phe	Gln 15	Leu	
Ser	Gln	Gly	Gly 20	Gln	Gly	Phe	Ala	Ile 25	Pro	Ile	Gly	Gln	Ala 30	Met	Ala	
Ile	Ala	Gly 35	Gln	Ile	Lys	Ĺeu	Pro	Thr	Val	His	Ile	Gly 45	Pro	Thr	Ala	
Phe	Leu 50	Gly	Leu	Gly	Val	Val 55	Asp	Asn	Asn	Gly	Asn 60	Gly	Ala	Arg	Val	
Gln 65		Val	Val	Gly	Ser 70		Pro	Ala	Ala	Ser 75		Gly	Ile	Ser	Thr 80	
	Asp	Val	Ile	Thr 85		Val	Asp	Gly	Ala 90	_	Ile	Asn	Ser	Ala 95		
Ala	Met	Ala	Asp 100	Ala	Leu	Asn	Gly	His 105	His	Pro	Gly	Asp	Val 110	Ile	Ser	
Val	Thr	Trp 115		Thr	Lys	Ser	Gly 120		Thr	Arg	Thr	Gly 125		Val	Thr	
Leu	Ala 130	Glu	Gly	Pro	Pro	Ala 135	Glu	Phe	Cys	Arg	Tyr 140	Pro	Ser	His	Trp	
Arg 145	Pro	Leu	Met	Phe	Ile 150	Thr	Lys	Glu	Leu	Met 155	Asn	Arg	Val	Ile	Glu 160	
Ile	His	Ala	His	Tyr 165	Asp	Gln	Arg	Gln	Leu 170	Ser	Gln	Ser	Pro	Asn 175	Thr	
Asn	Phe	Leu	Val 180	His	His	Pro	Tyr	Leu 185	Thr	Leu	Ile	Pro	Lys 190	Phe	Leu	
Leu	Gly	Ala 195	Leu	Ile	Val	Tyr	Ala 200	Pro	Tyr	Ser	Phe	Ala 205	Glu	Met	Glu	
Leu	Ala 210	Ile	Ser	Gly	His	Lys 215	Gln	Gly	Lys	Asp	Arg 220	Asp	Thr	Phe	Thr	
Met 225	Ile	Ser	Ser	Суѕ	Pro 230	Glu	Gly	Thr	Asn	Tyr 235	Ile	Ile	Asn	Arg	Lys 240	
Leu	Ile	Leu	Ser	Asp 245		Ser						Ser	Ser	Gly 255	-	
Ala	Phe	Arg	Asn 260	Leu	Ala	Gly	Lys	Ile 265	Ser	Phe	Leu	Gly	Lys 270	Asn	Ser	
Ser	Ala	Ser 275	Ile	His	Phe	Lys	His 280	Ile	Asn	Ile	Asn	Gly 285	Phe	Gly	Ala	
Gly	Vạl 290	Phe	Ser	Glu	Ser	Ser 295	Ile	Glu	Phe	Thr	Asp 300	Leu	Arg	Lys	Leu	
Val 305	Ala	Phe	Gly	Ser	Glu 310	Ser	Thr	Gly	Gly	Ile 315	Phe	Thr	Ala	Lys	Glu 320	
	Ile	Ser	Phe	Lys 325	Asn	Asn	His	His	11e 330	Ala	Phe	Arg	Asn	Asn 335		
Thr	Lys	Gly	Asn 340	Gly	Gly	Val	Ile	Gln 345		Gln	Gly	Asp	Met 350		Gly	
Ser	Val	Ser 355	Phe	Val	Asp	Gln	Arg 360		Ala	Ile	Ile	Phe 365		Asn	Asn	
Gln	Ala		Thr	Ser	Ser	Ser	Met	Lys	His	Ser	Gly	Arg	Gly	Gly	Ala	

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Ile	370 Ser	Glv	Asp	Phe	Ala	375 Glv	Ser	Arα	Tle	Leu	380 Phe	Leu	Asn	Asn	Gln		
385	001	OLy	пор	1,110	390	0-1	501	****		395	2.110	200			400		
Gln	Ile	Thr	Phe	Glu 405	Gly	Asn	Ser	Ala	Val 410	His	Gly	Gly	Ala	Ile 415	Tyr		
Asn	Lys	Asn	Gly 420	Leu	Val	Glu	Phe	Leu 425	Gly	Asn	Ala	Gly	Pro 430	Leu	Ala		
Phe	Lys	Glu 435	Asn	Thr	Thr	Ile	Ala 440		Gly	Gly	Ala	Ile 445		Thr	Ser		
Asn	Phe 450		Ala	Asn	Gln	Gln 455		Ser	Pro	Ile	Leu 460		Ser	Gln	Asn		
His 465		Asn	Lys	Lys	Gly 470		Ala	Ile	Tyr	Ala 475		Tyr	Val	Asn	Leu 480		
	Gln	Asn	Gln	Asp 485		Ile	Arg	Phe	Glu 490		Asn	Thr	Ala	Lys 495			
Gly	Gly	Gly	Ala 500	`	Thr	Ser	Ser	Gln 505		Ser	Ile	Thr	Ala 510		Asn		
Thr	Ile	Thr 515	Phe	Ser	Asp	Asn	Ala 520		Gly	Asp	Leu	Gly 525		Gly	Ala		
Ile	Leu 530		Glu	Gly	Lys	Lys 535		Ser	Leu	Thr	Leu 540		Ala	His	Ser		
Gly 545		Ile	Ala	Phe	Ser 550		Asn	Thr	Met	Leu 555		Ile	Thr	Lys	Lys 560		
	Ser	Leu	Asp	Arg 565		Asn	Ser	Ile	Leu 570		Lys	Glu	Ala	Pro 575			
Lys	Ile	Gln	Leu 580		Ala	Asn	Гуs	Asn 585		Ser	Ile	His	Phe 590		Asp		
Pro	Val	Met 595	Ala	Leu	Ser	Ala	Ser 600		Ser	Pro	Ile	Gln 605		Asn	Ala		
Pro	Glu 610		Glu	Thr	Pro	Phe 615		Ser	Pro	Lys		000				•	
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atgo cago acco ggco ggco gcgo	ggati gttca gcacg gacgi cttaa	tcg of the transfer of the tra	ccati tcggq tccaa tcaca ggcat	ceega geeta aegeg egegg	at co ac co gt go gt co cc co	gggca gcctt gtcgg gacgg ggtga	aggco ceto ggago geget aegto	g ato ggo gct gct cco cato	ggega ettgg ecegg gatea etegg	atcg ggtg gcgg act gtga	ttgt caag cggc	gccaq cgad gtctd ccacd ggcad	gat o caa o cgg o cgc o aac o	caago caaco catco gatgo caago	ggtggg cttccc ggcaac cccacc gcggac ccgggc	60 120 180 240 300 360 420	

135

480

540

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660

720 780

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1320 1380

1440

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1680

1740

1800

1860

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1980

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PCT/US01/23121

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Ser Gln Gly Gln Gly Phe Ala Ile Pro Ile Gly Gln Ala Met Ala Ile Ala Gly Gln Ile Lys Leu Pro Thr Val His Ile Gly Pro Thr Ala Phe Leu Gly Leu Gly Val Val Asp Asn Asn Gly Asn Gly Ala Arg Val 55 Gln Arg Val Val Gly Ser Ala Pro Ala Ala Ser Leu Gly Ile Ser Thr 70 75 Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr 90 Ala Met Ala Asp Ala Leu Asn Gly His His Pro Gly Asp Val Ile Ser 105 Val Thr Trp Gln Thr Lys Ser Gly Gly Thr Arg Thr Gly Asn Val Thr 115 120 125 Leu Ala Glu Gly Pro Pro Ala Glu Phe Cys Arg Tyr Pro Ser His Trp 135 Arg Pro Leu Met Pro Phe Ser Leu Arg Ser Thr Ser Phe Cys Phe Leu 155 150 Ala Cys Leu Cys Ser Tyr Ser Tyr Gly Phe Ala Ser Ser Pro Gln Val 165 170 175 Leu Thr Pro Asn Val Thr Thr Pro Phe Lys Gly Asp Asp Val Tyr Leu 185 Asn Gly Asp Cys Ala Phe Val Asn Val Tyr Ala Gly Ala Glu Asn Gly 200 Ser Ile Ile Ser Ala Asn Gly Asp Asn Leu Thr Ile Thr Gly Gln Asn 215 220 His Thr Leu Ser Phe Thr Asp Ser Gln Gly Pro Val Leu Gln Asn Tyr 230 235 Ala Phe Ile Ser Ala Gly Glu Thr Leu Thr Leu Lys Asp Phe Ser Ser 245 250 Leu Met Phe Ser Lys Asn Val Ser Cys Gly Glu Lys Gly Met Ile Ser 265 Gly Lys Thr Val Ser Ile Ser Gly Ala Gly Glu Val Ile Phe Trp Asp 280 Asn Ser Val Gly Tyr Ser Pro Leu Ser Ile Val Pro Ala Ser Thr Pro 295 300 Thr Pro Pro Ala Pro Ala Pro Ala Pro Ala Ala Ser Ser Leu Ser 310 Pro Thr Val Ser Asp Ala Arg Lys Gly Ser Ile Phe Ser Val Glu Thr 325 330 Ala Gly Asn Phe Gly Thr Val Phe Arg Gly Asn Ser Asn Asn Ala 360 Gly Ser Gly Ser Gly Ser Ala Thr Thr Pro Ser Phe Thr Val Lys 375 Asn Cys Lys Gly Lys Val Ser Phe Thr Asp Asn Val Ala Ser Cys Gly 390 395 Gly Gly Val Val Tyr Lys Gly Thr Val Leu Phe Lys Asp Asn Glu Gly 405 410 Gly Ile Phe Phe Arg Gly Asn Thr Ala Tyr Asp Asp Leu Gly Ile Leu 420 425 Ala Ala Thr Ser Arg Asp Gln Asn Thr Glu Thr Gly Gly Gly Gly 440 Val Ile Cys Ser Pro Asp Asp Ser Val Lys Phe Glu Gly Asn Lys Gly 455 460 Ser Ile Val Phe Asp Tyr Asn Phe Ala Lys Gly Arg Gly Ser Ile 470 475 Leu Thr Lys Glu Phe Ser Leu Val Ala Asp Asp Ser Val Val Phe Ser 490 Asn Asn Thr Ala Glu Lys Gly Gly Gly Ala Ile Tyr Ala Pro Thr Ile

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	Pro	Ser	Pro	Ser 165		Ser	Ser	Met	Pro 170		Ala	Val	Thr	Ile 175	
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Gly	Ala	Ser	Leu	Ser 245		Val	Ala	Gly	Ala 250		Asn	Asn	Asn	Tyr 255	
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Gln Arg Val Val Gly Ser Ala Pro Ala Ala Ser Leu Gly Ile Ser Thr
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Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr
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Ala Met Ala Asp Ala Leu Asn Gly His His Pro Gly Asp Val Ile Ser
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Ser Thr Ile Ser Leu Ser Gly Ile Thr Lys Ala Thr Phe Ser Cys Asn
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Ile Tyr Ala Lys Gly Asp Leu Thr Ile Ala Asp Ser Gln Glu Val Leu
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Phe Ser Ile Asn Lys Ala Thr Lys Asp Gly Gly Ala Ile Phe Ala Glu
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Lys Asp Val Ser Phe Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn 450 460
Gly Ala Glu Glu Lys Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser 465 470 475 480
Ile Gln Ser Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln
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Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr
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151

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WO 02/08267 PCT/US01/23121

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Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr
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70

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160

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162

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Gln Ala Thr Ala Tyr Asp Thr Leu Asp Ile Asn Lys Ser Glu Asp Ser
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Gly Ser Leu Val Leu Lys Pro Asn Thr Glu Leu His Val Ile Ser Phe
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WO 02/08267

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Arg Pro Glu Asn Pro Phe Leu Ser Ser Gly Pro Phe Met Pro Lys Thr
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Ser Ala Leu Tyr Thr His Tyr Pro Val Asp Gly Thr Phe Trp Leu Ile
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WO 02/08267

166

PCT/US01/23121

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WO 02/08267

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179

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Arg	Gln 290	Gly	Gly	Thr	His	Leu 295	Thr	Gly	Phe	Ser	Thr 300	Ala	Leu	Thr	Arg
Val 305	Ile	Asn	Thr	Tyr	Ile 310	Lys	Ala	His	Asn	Leu 315	Ala	Lys	Asn	Asn	Lys 320
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Ala	Leu 370	Thr	Ile	Phe	Phe	Glu 375	Glu	Asn	Pro	Gln	Ile 380	Ala	Arg	Met	Ile
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Gly	Thr	Ile	Ile	Ala 485	Ala	Leu	Gly	Суз	Gly 490	Ile	Gly	Ala	Asp	Asn 495	Phe
Asn	Leu	Ser	Lys 500	Leu	Arg	Tyr	Arg	Arg 505	Ile	Ile	Ile	Met	Thr 510	Asp	Ala
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181

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182

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Ile	Glu	Asn	Ala	Ala 245	Lys	Arg	Gly	Thr	Ile 250	Ļys	Ile	Asp	Thr	Ile 255	Gln
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Ser	Arg	Ala 275	Lys	Glu	Met	Leu	Pro 280	Leu	Leu	Phe	Glu	His 285	Thr	Glu	Суз
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Gln	Gly	Tyr	Leu	Glu 325	Lys	Glu	Leu	Leu	Leu 330	Leu	Gln	Glu	Gln	Leu 335	Thr
	-		Tyr 340		_			345	_				350		-
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His 385	Glu	Leu	Ala	Thr	Pro 390	Val	Thr	Lys	Gln	Asp 395	Thr	Ser	Gln	Leu	Ala 400
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-			Leu 420				_	425					430	_	-
		435	Ile				440					445			
	450		Gly			455					460			Asn	Phe
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Leu

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<212> PRT

<213> Chlamydia pneumoniae

<400> 391

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20 25 30

Ser Ala Ser Asp Ala Ile Lys Lys Leu Trp Tyr Leu Glu Leu Lys Asp

WO 02/08267 PCT/US01/23121

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Lys Val Leu Gly Pro Arg Asn Leu Met Pro Thr Pro Lys Thr Gly Thr 130 135 140

Val Thr Thr Asp Val Ala Lys Ala Ile Ser Glu Leu Arg Lys Gly Lys 145 150 155 160

Ile Glu Phe Lys Ala Asp Arg Ala Gly Val Cys Asn Val Gly Val Gly 165. 170 175

Lys Leu Ser Phe Glu Ser Ser Gln Ile Lys Glu Asn Ile Glu Ala Leu 180 185 190

Ser Ser Ala Leu Ile Lys Ala Lys Pro Pro Ala Ala Lys Gly Gln Tyr 195 200 205

Leu Val Ser Phe Thr Ile Ser Ser Thr Met Gly Pro Gly Ile Ser Ile 210 220

Asp Thr Arg Glu Leu Met Ala Ser 225 230

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. <211> 122

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<213> Chlamydia pneumoniae

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Ile Ile Lys Lys Leu Lys Leu Asp Pro Glu Ala Arg Ala Ser Glu Leu 35 40 45

Thr Glu Glu Glu Val Gly Arg Leu Asn Ser Leu Leu Gln Ser Glu Tyr 50 60

Thr Val Glu Gly Asp Leu Arg Arg Val Gln Ser Asp Ile Lys Arg 65 70 75 80

Leu Ile Ala Ile His Ser Tyr Arg Gly Gln Arg His Arg Leu Ser Leu 85 90 95

Pro Val Arg Gly Gln Arg Thr Lys Thr Asn Ser Arg Thr Arg Lys Gly 100 105 110

Lys Arg Lys Thr Val Ala Gly Lys Lys 115 120

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<211> 1723

<212> PRT

<213> Chlamydia pneumoniae

<400> 394

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Thr	Ile	Asp	Thr 340	Ser	Thr	Leu	Gln	Thr 345	Arg	Ala	Ala	Ser	Ala 350	Thr	Pro
Ala	Val	Ala 355	Pro	Val	Ala	Ala	Val 360	Thr	Pro	Thr _.	Pro	Ile 365	Ser	Thr	Gln
Glu	Thr 370	Ala	Gly	Asn	Gly	Gly 375	Ala	Ile	Tyr	Ala	Lys 380	Gln	Gly	Ile	Ser
Ile 385	Ser	Thr	Phe	Lys	Asp 390	Leu	Thr	Phe	Lys	Ser 395	Asn	Ser	Ala	Ser	Val 400
Asp	Ala	Thr	Leu	Thr 405	Val	Asp	Ser	Ser	Thr 410	Ile	Gly	Glu	Ser	Gly 415	Gly
Ala	Ile	Phe	Ala 420	Ala	Asp	Ser	Ile	Gln 425	Ile	Gln	Gln	Суѕ	Thr 430	Gly	Thr
Thr	Leu	Phe 435	Ser	Gly	Asn	Thr	Ala 440	Asn	Lys	Ser	Gly	Gly 445	Gly	Ile	Tyr
Ala	Val 450	СſУ	Gln	Val	Thr	Leu 455	Glu	Asp	Ile	Ala	Asn 460	Leu	Lys	Met	Thr .
Asn 465	Asn	Thr	Cys	Lys	Gly 470	Glu	Gly	Gly	Ala	Ile 475	Tyr	Thr	Lys	Lys	Ala 480
Leu	Thr	Ile	Asn	Asn 485	Gly	Ala	Ile	Leu	Thr 490	Thr	Phe	Ser	Gly	Asn 495	Thr
Ser	Thr	Asp	Asn 500	Gly	Gly	Ala	Ile	Phe 505	Ala	Val	Gly	Gly	Ile 510	Thr	Leu
Ser	Asp	Leu 515	Val	Glu	Val	Arg	Phe 520	Ser	Lys	Asn	Lys	Thr 525	Gly	Asn	Tyr
Ser	Ala 530		Ile	Thr	Lys	Ala 535	Ala	Ser	Asn	Thr	Ala 540	Pro	Val	Val ·	Ser
Ser 545	Ser	Thr	Thr	Ala	Ala 550	Ser	Pro	Ala	Val	Pro 555	Ala	Ala	Ala	Ala	Ala 560
Pro	Val	Thr	Asn	Ala 565	Ala	Lys	Gly	Gly	Ala 570	Leu	Tyr	Ser	Thr	Glu 575	Gly
Leu	Thr	Val	Ser 580	Gly	Ile	Thr	Ser	Ile 585	Leu	Ser	Phe	Glu	Asn 590	Asn	Glu
Cys	Gln	Asn 595		Gly	Gly	Gly	Ala 600	Tyr	Val	Thr	Lys	Thr 605	Phe	Gln	Cys
Ser	Asp 610		His	Arg	Leu	Gln 615	Phe	Thr	Ser	Asn	Lys 620	Ala	Ala	Asp	Glu
Gly 625		Gly	Leu	Tyr	Cys 630	Gly	Asp	Asp	Val	Thr 635	Leu	Thr	Ası	Leu	Thr 640
Gly	Lys	Thr	Leu	Phe 645	Gln	Glu	Asn	Ser	Ser 650	Glu	Lys	His	Gly	Gly 655	Gly

PCT/US01/23121 WO 02/08267

Leu	Ser	Leu	Ala 660	Ser	Gly	Lys	Ser	Leu 665	Thr	Met	Thr	Ser	Leu 670	Glu	Ser
Phe	Cys	Leu 675	Asn	Ala	Asn	Thr	Ala 680	Lys	Glu	Asn	Glÿ	Gly 685	Gly	Ala	Asn
Val	Pro 690	Glu	Asn	Ile	Val	Leu 695	Thr	Phe	Thr	Tyr	Thr 700	Pro	Thr	Pro	Asn
Glu 705	Pro	Ala	Pro	Val	Gln 710	Gln	Pro	Val	Tyr	Gly 715	Glu	Ala	Leu	Val	Thr 720
Gly	Asn	Thr	Ala	Thr 725	Lys	Ser	Gly	Gly	Gly 730	Ile	Tyr	Thr	Lys	Asn 735	Ala
Ala	Phe	Ser	Asn 740	Leu	Ser	Ser	Val	Thr 745	Phe	Asp	Gln	Asn	Thr 750	Ser	Ser
Glu	Asn	Gly 755	Gly	Ala	Leu	Leu	Thr 760	Gln	Lys	Ala	Ala	Asp 765	Lys	Thr	Asp
Cys	Ser 770	Phe	Thr	Tyr	Ile	Thr 775	Asn	Val	Asn	Ile	Thr 780	Asn	Asn	Thr	Ala
Thr 785	Gly	Asn	Gly	Gly	Gly 790	Ile	Ala	Gly	Gly	Lys 795	Ala	His	Phe	Asp	Arg 800
Ile	Asp	Asn	Leu	Thr 805	Val	Gln	Ser	Asn	Gln 810	Ala	Lys	Lys	Gly	Gly 815	Gly
Val	Tyr	Leu	Glu 820	Asp	Ala	Leu	Ile	Leu 825	Glu	Lys	Val	Ile	Thr 830	Gly	Ser
Val	Ser	Gln 835		Thr	Ala	Thr	Glu 840	Ser	Gly	Gly	Gly	Ile 845	Tyr	Ala	Lys
Asp	Ile 850	Gln	Leu	Gln	Ala	Leu 855	Pro	Gly	Ser	Phe	Thr 860	Ile	Thr	Asp	Asn
865					870					875				Gly	880
				885					890		•	•		Thr 895	
			900					905					710		
_		915					920					925	•	lle	
Gln	Cys 930		Thr	Pro	Ile	Leu 935	Phe	Ser	Asn	Asn	Ser 940	Ala	Ala	Thr	Lys
945					950					955	ı			e Ser	900
				965					970					975	
Ser	Ala	Lys	Ser	Glu	Ala	Thr	Thr	Ala	Ala	Thr	Ala	Gl	, Asr	ı Lys	Asp

990 985 980 Ser Cys Gly Gly Ala Ile Ala Ala Asn Ser Val Thr Leu Thr Asn Asn 1000 995 Pro Glu Ile Thr Phe Lys Gly Asn Tyr Ala Glu Thr Gly Gly Ala Ile 1015 1020 Gly Cys Ile Asp Leu Thr Asn Gly Ser Pro Pro Arg Lys Val Ser Ile Ala Asp Asn Gly Ser Val Leu Phe Gln Asp Asn Ser Ala Leu Asn Arg 1050 1045 Gly Gly Ala Ile Tyr Gly Glu Thr Ile Asp Ile Ser Arg Thr Gly Ala 1065 Thr Phe Ile Gly Asn Ser Ser Lys His Asp Gly Ser Ala Ile Cys Cys 1080 Ser Thr Ala Leu Thr Leu Ala Pro Asn Ser Gln Leu Ile Phe Glu Asn 1095 Asn Lys Val Thr Glu Thr Thr Ala Thr Thr Lys Ala Ser Ile Asn Asn 1110 Leu Gly Ala Ala Ile Tyr Gly Asn Asn Glu Thr Ser Asp Val Thr Ile 1130 1125 Ser Leu Ser Ala Glu Asn Gly Ser Ile Phe Phe Lys Asn Asn Leu Cys 1145 Thr Ala Thr Asn Lys Tyr Cys Ser Ile Ala Gly Asn Val Lys Phe Thr 1160 Ala Ile Glu Ala Ser Ala Gly Lys Ala Ile Ser Phe Tyr Asp Ala Val 1175 Asn Val Ser Thr Lys Glu Thr Asn Ala Gln Glu Leu Lys Leu Asn Glu 1195 1190 Lys Ala Thr Ser Thr Gly Thr Ile Leu Phe Ser Gly Glu Leu His Glu Asn Lys Ser Tyr Ile Pro Gln Lys Val Thr Phe Ala His Gly Asn Leu 1220 Ile Leu Gly Lys Asn Ala Glu Leu Ser Val Val Ser Phe Thr Gln Ser 1245 1240 Pro Gly Thr Thr Ile Thr Met Gly Pro Gly Ser Val Leu Ser Asn His Ser Lys Glu Ala Gly Gly Ile Ala Ile Asn Asn Val Ile Ile Asp Phe 1270 1275

Ser Glu Ile Val Pro Thr Lys Asp Asn Ala Thr Val Ala Pro Pro Thr

Leu Lys Leu Val Ser Arg Thr Asn Ala Asp Ser Lys Asp Lys Ile Asp 1300 1305 1310

- Ile Thr Gly Thr Val Thr Leu Leu Asp Pro Asn Gly Asn Leu Tyr Gln 1320
- Asn Ser Tyr Leu Gly Glu Asp Arg Asp Ile Thr Leu Phe Asn Ile Asp 1340 1335
- Asn Ser Ala Ser Gly Ala Val Thr Ala Thr Asn Val Thr Leu Gln Gly 1350
- Asn Leu Gly Ala Lys Lys Gly Tyr Leu Gly Thr Trp Asn Leu Asp Pro 1370
- Asn Ser Ser Gly Ser Lys Ile Ile Leu Lys Trp Thr Phe Asp Lys Tyr 1380
- Leu Arg Trp Pro Tyr Ile Pro Arg Asp Asn His Phe Tyr Ile Asn Ser
- Ile Trp Gly Ala Gln Asn Ser Leu Val Thr Val Lys Gln Gly Ile Leu
- Gly Asn Met Leu Asn Asn Ala Arg Phe Glu Asp Pro Ala Phe Asn Asn 1435 1430
- Phe Trp Ala Ser Ala Ile Gly Ser Phe Leu Arg Lys Glu Val Ser Arg 1450
- Asn Ser Asp Ser Phe Thr Tyr His Gly Arg Gly Tyr Thr Ala Ala Val
- Asp Ala Lys Pro Arg Gln Glu Phe Ile Leu Gly Ala Ala Phe Ser Gln
- Val Phe Gly His Ala Glu Ser Glu Tyr His Leu Asp Asn Tyr Lys His 1495
- Lys Gly Ser Gly His Ser Thr Gln Ala Ser Leu Tyr Ala Gly Asn Ile 1515
- Phe Tyr Phe Pro Ala Ile Arg Ser Arg Pro Ile Leu Phe Gln Gly Val 1530
- Ala Thr Tyr Gly Tyr Met Gln His Asp Thr Thr Thr Tyr Tyr Pro Ser 1545
- Ile Glu Glu Lys Asn Met Ala Asn Trp Asp Ser Ile Ala Trp Leu Phe 1555
- Asp Leu Arg Phe Ser Val Asp Leu Lys Glu Pro Gln Pro His Ser Thr 1575
- Ala Arg Leu Thr Phe Tyr Thr Glu Ala Glu Tyr Thr Arg Ile Arg Gln 1595 1590
- Glu Lys Phe Thr Glu Leu Asp Tyr Asp Pro Arg Ser Phe Ser Ala Cys 1610 1605
- Ser Tyr Gly Asn Leu Ala Ile Pro Thr Gly Phe Ser Val Asp Gly Ala 1625

Leu Ala Trp Arg Glu Ile Ile Leu Tyr Asn Lys Val Ser Ala Ala Tyr 1635 1640 1645

Leu Pro Val Ile Leu Arg Asn Asn Pro Lys Ala Thr Tyr Glu Val Leu 1650 1655 1660

Ser Thr Lys Glu Lys Gly Asn Val Val Asn Val Leu Pro Thr Arg Asn 1665 1670 1675 1680

Ala Ala Arg Ala Glu Val Ser Ser Gln Ile Tyr Leu Gly Ser Tyr Trp 1685 1690 1695

Thr Leu Tyr Gly Thr Tyr Thr Ile Asp Ala Ser Met Asn Thr Leu Val 1700 1705 1710

Gln Met Ala Asn Gly Gly Ile Arg Phe Val Phe 1715 1720

<210> 395

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<212> PRT

<213> Chlamydia pneumoniae

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305	Lys				310					315				Ala	320
Ala				325					330					Ser 335	
			340	Ser				345					350	Thr	
		355					360					365		Thr	
	370					375					380			Ile	
205					390					395				Ser	400
				405					410					Gly 415	•
			420					425					430	Gly	
		435					440					445		Ile	
	450	_				455					. 460			Met	
465					470					4/5				Lys	400
Leu				485					490					Asn 495	
Ser			500					505					210	Thr	
	-	515					520					525		Asn Val	
	530					535					540			Ala	
545					550					555				Glu	260
				565					570					575 Asn	
			580					585					590	Gln	
_		595					600					605		Asp	
	610					615					620			Leu	
625					630					635				Gly	640
_				645					650					655 Glu	
			660					665					6/0	Ala	
		675					680					685	ı	Pro	
•	600					695					700			. Val	
705					710					715				Asn	120
				725					/30					735 Ser	
			740					/45					/50) Thr	
		755					760					/65)	1 Thr	
~ys	عرجاد			3											

	770					775					780			_	
Thr 785	Gly	Asn	Gly	Gly	Gly 790	Ile	Ala	Gly	Gly	Lys 795	Ala	His	Phe	Asp	Arg 800
Ile	Asp	Asn	Leu	Thr 805	Val	Gln	Ser	Asn	Gln 810	Ala	Lys	Lys	Gly	Gly 815	Gly
Val	Tyr	Leu	Glu 820		Ala	Leu	Ile	Leu 825		Lys	Val	Ile	Thr 830	Gly	Ser
Val	Ser	Gln 835		Thr	Ala	Thr	Glu 840		Gly	Gly	Gly	Ile 845	Tyr	Ala	ГЛЗ
Asp	Ile 850		Leu	Gln	Ala	Leu 855		Gly	Ser	Phe	Thr 860	-	Thr	Asp	Asn
Lys 865	Val	Glu	Thr	Ser	Leu 870		Thr	Ser	Thr	Asn 875		Tyr	Gly	Gly	Gly 880
Ile	Tyr	Ser	Ser	Gly 885		Val	Thr	Leu	Thr 890		Ile	Ser	Gly	Thr 895	Phe
Gly	Ile	Thr	Gly 900		Ser	Val	Ile	Asn 905		Ala	Thr	Ser	Gln 910	Asp	Ala
Asp	Ile	Gln 915		Gly	Gly	Ile	Tyr 920		Thr	Thr	Ser	Leu 925	Ser	Ile	Asn
	Cys 930	Asn				935			•		940				
945					950					955					960
				965					970					975	Asn ·
	Ala	_	980					985					990		
	Cys	995					1000)				100	5		
	Glu 101	0				1015	5				1020)			
102					1030)				1039	5				1040
	Asp			1045	5				1050)				105	5
_	Gly		1060)				1065	5				107	0	
Thr			~ 7	Asn	Ser	802	T			$C1$ α	Ser	Ala	TIE	Cys	Cys
	Phe	107	5				1080)				108	5		
	Thr	107: Ala 0	5 Leu	Thr	Leu	Ala 1095	1080 Pro) Asn	Ser	Gln	Leu 1100	108: Ile)	5 Phe	Glu	Asn
Asr.	Thr 109 Lys	107: Ala O Val	5 Leu Thr	Thr Glu	Leu Thr 1110	Ala 1095 Thr	1080 Pro Ala) Asn Thr	Ser Thr	Gln Lys 111	Leu 1100 Ala	108 Ile) Ser	5 Phe Ile	Glu Asn	Asn Asn 1120
Asr 110 Leu	Thr 109 Lys 5 Gly	107: Ala O Val Ala	5 Leu Thr Ala	Thr Glu Ile 1125	Leu Thr 1110 Tyr	Ala 1095 Thr) Gly	1080 Pro Ala Asn	Asn Thr Asn	Ser Thr Glu 1130	Gln Lys 111: Thr	Leu 1100 Ala Ser	108 Ile) Ser Asp	Fhe Ile Ile	Glu Asn Thr 113	Asn Asn 1120 Ile
Asr 110 Lev	Thr 109 Lys 5 Gly	107: Ala 0 Val Ala Ser	5 Leu Thr Ala Ala 1140	Thr Glu Ile 1125 Glu	Leu Thr 1110 Tyr Asn	Ala 1099 Thr Gly Gly	1080 Pro Ala Asn Ser	Asn Thr Asn Ile	Ser Thr Glu 1130 Phe	Gln Lys 111: Thr) Phe	Leu 1100 Ala Ser Lys	108 Ile Ser Asp	Phe Ile Ile Asn 115	Glu Asn Thr 113 Leu 0	Asn Asn 1120 Ile 5 Cys
Asr 110 Lev Ser Thr	Thr 109 Lys 5 Gly Leu	107: Ala 0 Val Ala Ser Thr	5 Leu Thr Ala Ala 1140 Asn	Thr Glu Ile 1125 Glu) Lys	Leu Thr 1110 Tyr S Asn	Ala 1095 Thr) Gly Gly Cys	1080 Pro Ala Asn Ser Ser	Thr Asn Ile 1145 Ile	Ser Thr Glu 1130 Phe 5	Gln Lys 111! Thr) Phe Gly	Leu 1100 Ala Ser Lys Asn	108: Ile Ser Asp Asn Val	Phe Ile Ile Asn 115 Lys	Glu Asn Thr 113 Leu O Phe	Asn Asn 1120 Ile Cys Thr
Asr 110 Leu Ser Thr	Thr 109 Lys 5 Gly Leu Ala Ile	107: Ala 0 Val Ala Ser Thr 115: Glu	5 Leu Thr Ala Ala 1140 Asn 5 Ala	Thr Glu Ile 1125 Glu Lys Ser	Leu Thr 1110 Tyr Asn Tyr	Ala 1095 Thr Cly Cys Cys	1080 Pro Ala Asn Ser Ser 1160 Lys	Asn Thr Asn Ile 1145 Ile)	Ser Thr Glu 1130 Phe Ala Ile	Gln Lys 111: Thr Phe Gly Ser	Leu 1100 Ala Ser Lys Asn Phe 1180	108: Ile Ser Asp Asn Val 116: Tyr	Phe Ile Ile Asn 115 Lys Asp	Glu Asn Thr 113 Leu O Phe Ala	Asn Asn 1120 Ile 5 Cys Thr
Asn 110 Leu Ser Thr Ala	Thr 109 Lys 5 Gly Leu Ala 11e 117	1073 Ala O Val Ala Ser Thr 1153 Glu O Ser	5 Leu Thr Ala Ala 1140 Asn 5 Ala	Thr Glu Ile 1125 Glu Lys Ser Lys	Leu Thr 1110 Tyr Asn Tyr Ala Glu 1190	Ala 1095 Thr Gly Gly Cys Gly 1175 Thr	Ala Asn Ser Ser 1160 Lys Asn	Asn Thr Asn Ile 1145 Ile Ala Ala	Ser Thr Glu 1130 Phe 5 Ala Ile Gln	Gln Lys 1119 Thr Phe Gly Ser Glu 1199	Leu 1100 Ala Ser Lys Asn Phe 1180 Leu	IOS: Ile Ser Asp Asn Val 116 Tyr Uys	Phe Ile Ile Asn 115 Lys Asp	Asn Thr 113 Leu O Phe Ala Asn	Asn 1120 11e 5 Cys Thr Val Glu 1200
Asr 110 Leu Ser Thr Ala Asr 118 Lys	Thr 109 Lys Gly Leu Ala Ile 117 Val	1075 Ala O Val Ala Ser Thr 1155 Glu O Ser Thr	Thr Ala Ala 1140 Asn 5 Ala Thr	Thr Glu Ile 1125 Glu Lys Ser Lys Thr 1205	Thr 1110 Tyr Asn Tyr Ala Glu 1190 Gly	Ala 1095 Thr Gly Gly Cys Gly 1175 Thr	Ala Asn Ser Lys Asn Ile	Asn Thr Asn Ile 1145 Ile Ala Ala Leu	Ser Thr Glu 1130 Phe Ala Ile Gln Phe 1210	Gln Lys 1111 Thr Phe Gly Ser Glu 1199 Ser	Leu 1100 Ala Ser Lys Asn Phe 1180 Leu 5	108: Ile Ser Asp Asn Val 116: Tyr Lys	Phe Ile Ile Asn 115 Lys Asp Leu Leu	Glu Asn Thr 113 Leu O Phe Ala Asn His 121	Asn 1120 11e 5 Cys Thr Val Glu 1200 Glu 5
Asn 110 Ser Thr Ala Asn 118 Lys	Thr 109 Lys Gly Leu Ala Ile 117 Val	1079 Ala O Val Ala Ser Thr 1159 Glu O Ser Thr	Thr Ala Ala 1140 Asn Ala Thr Ser	Thr Glu Ile 1125 Glu Lys Ser Lys Thr 1205 Ile	Thr 1110 Tyr Asn Tyr Ala Glu 1190 Gly	Ala 1095 Thr Gly Gly Cys Gly 1175 Thr	Ala Asn Ser Ser Lys Asn Ile	Asn Thr Asn Ile 1145 Ile Ala Ala Leu Val 1225	Ser Thr Glu 1130 Phe 5 Ala Ile Gln Phe 1210 Thr	Gln Lys 111: Thr Phe Gly Ser Glu 119: Ser	Leu 1100 Ala Ser Lys Asn Phe 1180 Leu Gly	108: Ile Ser Asp Asn Val 116: Tyr Lys Glu	Phe Ile Ile Asn 115 Lys Asp Leu Gly 123	Glu Asn Thr 113 Leu O Phe Ala Asn His 121 Asn O	Asn 1120 11e 5 Cys Thr Val Glu 1200 Glu 5 Leu
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Ser Lys Glu Ala Gly Gly Ile Ala Ile Asn Asn Val Ile Ile Asp Phe 1270 1275 Ser Glu Ile Val Pro Thr Lys Asp Asn Ala Thr Val Ala Pro Pro Thr . 1290 1285 Leu Lys Leu Val Ser Arg Thr Asn Ala Asp Ser Lys Asp Lys Ile Asp 1305 1310 1300 Ile Thr Gly Thr Val Thr Leu Leu Asp Pro Asn Gly Asn Leu Tyr Gln 1315 1320 1325 Asn Ser Tyr Leu Gly Glu Asp Arg Asp Ile Thr Leu Phe Asn Ile Asp 1335 1340 Asn Ser Ala Ser Gly Ala Val Thr Ala Thr Asn Val Thr Leu Gln Gly 1350 1355 Asn Leu Gly Ala Lys Lys Gly Tyr Leu Gly Thr Trp Asn Leu Asp Pro 1365 1370 1375 Asn Ser Ser Gly Ser Lys Ile Ile Leu Lys Trp Thr Phe Asp Lys Tyr 1385 1390 1380 Leu Arg Trp Pro Tyr Ile Pro Arg Asp Asn His Phe Tyr Ile Asn Ser 1395 1400 1405 Ile Trp Gly Ala Gln Asn Ser Leu Val Thr Val Lys Gln Gly Ile Leu 1410 1415 1420 Gly Asn Met Leu Asn Asn Ala Arg Phe Glu Asp Pro Ala Phe Asn Asn 1425 1430 1435 Phe Trp Ala Ser Ala Ile Gly Ser Phe Leu Arg Lys Glu Val Ser Arg 1445 1450 Asn Ser Asp Ser Phe Thr Tyr His Gly Arg Gly Tyr Thr Ala Ala Val 1470 1465 1460 Asp Ala Lys Pro Arg Gln Glu Phe Ile Leu Gly Ala Ala Phe Ser Gln 1475 1480 1485 Val Phe Gly His Ala Glu Ser Glu Tyr His Leu Asp Asn Tyr Lys His 1490 1495 1500 . Lys Gly Ser Gly His Ser Thr Gln Ala Ser Leu Tyr Ala Gly Asn Ile 1510 1515 Phe Tyr Phe Pro Ala Ile Arg Ser Arg Pro Ile Leu Phe Gln Gly Val 1525 1530 1535 Ala Thr Tyr Gly Tyr Met Gln His Asp Thr Thr Tyr Tyr Pro Ser 1540 1545 1550 1540 Ile Glu Glu Lys Asn Met Ala Asn Trp Asp Ser Ile Ala Trp Leu Phe 1555 1560 1565 Asp Leu Arg Phe Ser Val Asp Leu Lys Glu Pro Gln Pro His Ser Thr 1570 1575 1580 Ala Arg Leu Thr Phe Tyr Thr Glu Ala Glu Tyr Thr Arg Ile Arg Gln 1590 1595 Glu Lys Phe Thr Glu Leu Asp Tyr Asp Pro Arg Ser Phe Ser Ala Cys 1610 1615 1605 Ser Tyr Gly Asn Leu Ala Ile Pro Thr Gly Phe Ser Val Asp Gly Ala 1620 1625 1630 Leu Ala Trp Arg Glu Ile Ile Leu Tyr Asn Lys Val Ser Ala Ala Tyr 1635 1640 1645 Leu Pro Val Ile Leu Arg Asn Asn Pro Lys Ala Thr Tyr Glu Val Leu 1660 1650 1655 Ser Thr Lys Glu Lys Gly Asn Val Val Asn Val Leu Pro Thr Arg Asn 1665 1670 1675 168 Ala Ala Arg Ala Glu Val Ser Ser Gln Ile Tyr Leu Gly Ser Tyr Trp 1685 1690 Thr Leu Tyr Gly Thr Tyr Thr Ile Asp Ala Ser Met Asn Thr Leu Val 1700 1705 Gln Met Ala Asn Gly Gly Ile Arg Phe Val Phe 1720

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<400> 396

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Glu Glu Val Phe Arg Glu Ile Phe Pro Ile Lys Ser Tyr Asn Glu Ala 50 55 60

Thr Val Leu Glu Tyr Leu Ser Tyr Asn Leu Gly Val Pro Lys Tyr Ser 65 70 75 80

Pro Glu Glu Cys Ile Arg Gly Ile Thr Tyr Ser Val Thr Leu Lys 85 90 95

Val Arg Phe Arg Leu Thr Asp Glu Thr Gly Ile Lys Glu Glu Val 100 105 110

Tyr Met Gly Thr Ile Pro Leu Met Thr Asp Lys Gly Thr Phe Ile Ile 115 120 125

Asn Gly Ala Glu Arg Val Val Val Ser Gln Val His Arg Ser Pro Gly 130 135 140

Ile Asn Phe Glu Gln Glu Lys His Ser Lys Gly Asn Ile Leu Phe Ser 145 150 155 160

Phe Arg Ile Ile Pro Tyr Arg Gly Ser Trp Leu Glu Ala Ile Phe Asp 165 170 175

Ile Asn Asp Leu Ile Tyr Ile His Ile Asp Arg Lys Lys Arg Arg Arg 180 185 . 190

Lys Ile Leu Ala Ile Thr Phe Ile Arg Ala Leu Gly Tyr Ser Ser Asp 195 200 205

Ala Asp Ile Ile Glu Glu Phe Phe Thr Ile Gly Glu Ser Ser Leu Arg 210 215 220

Ser Glu Lys Asp Phe Ala Leu Leu Val Gly Arg Ile Leu Ala Asp Asn 225 230 235 240

Ile Ile Asp Glu Ala Ser Ser Leu Val Tyr Gly Lys Ala Gly Glu Lys 245 250 255

Leu Ser Thr Ala Met Leu Lys Arg Met Leu Asp Ala Gly Ile Ala Ser 260 265 270

Val Lys Ile Ala Val Asp Ala Asp Glu Asn His Pro Ile Ile Lys Met 275 280 285

Leu Ala Lys Asp Pro Thr Asp Ser Tyr Glu Ala Ala Leu Lys Asp Phe 290 295 300

Tyr 305	Arg	Arg	Leu	Arg	Pro 310	Gly	Glu	Pro	Ala	Thr 315	Leu	Ala	Asn	Ala	Arg 320
Ser	Thr	Ile	Met	Arg 325	Leu	Phe	Phe	Asp	Pro 330	Lys	Arg	Tyr	Asn	Leu 335	Gly
Arg	Val	Gly	Arg 340	Tyr	Lys	Leu	Asn	Arg 345	Lys	Leu	Gly	Phe	Ser 350	Ile	Asp
Asp	Glu	Ala 355	Leu	Ser	Gln	Val	Thr 360	Leu	Arg	Lys	Glu	Asp 365	Val	Ile	Gly
Ala	Leu 370	Lys	Tyr	Leu	Ile	Arg 375	Leu	Lys	Met	Gly	Asp 380	Glu	Lys	Ala	Cys ·
Val 385	Asp	Asp	Ile	Asp	His 390	Leu	Ala	Asn	Arg	Arg 395	Val	Arg	Ser	Val	Gly 400
Glu	Leu	Ile	Gln	Asn 405	Gln	Cys	Arg	Ser	Gly 410	Leu	Ala	Arg	Met	Glu 415	Lys
Ile	Val	Arg	Glu 420	Arg	Met	Asn	Leu	Phe 425	Asp	Phe	Ser	Ser	Asp 430	Thr	Leu
Thr	Pro	Gly 435	Lys	Val	Val	Ser	Ala 440	Lys	Gly	Leu	Ala	Ser 445	Val	Leu	Lys
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Pro 465	Val	Ala	Glu	Leu	Thr 470	His	Lys	Arg	Arg	Leu 475	Ser	Ala	Leu	Gly	Pro 480
Gly	Gly	Leu	Asn	Arg 485	Glu	Arg	Ala	Gly	Phe 490	Glu	Val	Arg	Asp	Val 495	His
Ala	Ser	His	Tyr 500	Gly	Arg	Ile	Cys	Pro 505	Ile	Glu	Thr	Pro	Glu 510	Gly	Pro
Asn	Ile	Gly 515	Leu	Ile	Thr	Ser	Leu 520	Ser	Ser	Phe	Ala	Lys 525	Ile	Asn	Glu
Phe	Gly 530	Phe	Ile	Glu	Thr	Pro 535	Tyr	Arg	Ile	Val	Arg 540	Asp	Gly	Ile	Val
Thr 545	Asp	Glu	Ile	Glu	Tyr 550	Met	Thr	Ala	Asp	Val 555	Glu	Glu	Glu	Сув	Val 560
Ile	Ala	Gln	Ala	Ser 565	Ala	Ser	Leu	Asp	Glu 570	Tyr	Asn	Met	Phe	Thr 575	Glu
Pro	Val	Суѕ	Trp 580	Val	Arg	Tyr	Ala	Gly 585	Glu	Ala	Phe	Glu	Ala 590	Asp	Thr
Ser	Thr	Val 595	Thr	His	Met	Asp	Val 600	Ser	Pro	Lys	Gln	Leu 605	Val	Ser	Ile
Val	Thr 610	Gly	Leu	Ile	Pro	Phe 615	Leu	Glu	His	Asp	Asp 620	Ala	Asņ	Arg	Ala

Leu Met Gly Ser Asn Met Gln Arg Gln Ala Val Pro Leu Leu Lys Thr 630 625 635 Glu Ala Pro Val Val Gly Thr Gly Leu Glu Cys Arg Ala Ala Lys Asp 650 Ser Gly Ala Ile Val Val Ala Glu Glu Asp Gly Val Val Asp Phe Val 665 Asp Gly Tyr Lys Val Val Val Ala Ala Lys His Asn Pro Thr Ile Lys 680 Arg Thr Tyr His Leu Lys Lys Phe Leu Arg Ser Asn Ser Gly Thr Cys Ile Asn Gln Gln Pro Leu Cys Ala Val Gly Asp Val Ile Thr Lys Gly Asp Val Ile Ala Asp Gly Pro Ala Thr Asp Arg Gly Glu Leu Ala Leu Gly Lys Asn Val Leu Val Ala Phe Met Pro Trp Tyr Gly Tyr Asn Phe Glu Asp Ala Ile Ile Ile Ser Glu Lys Leu Ile Arg Glu Asp Ala Tyr Thr Ser Ile Tyr Ile Glu Glu Phe Glu Leu Thr Ala Arg Asp Thr Lys Leu Gly Lys Glu Glu Ile Thr Arg Asp Ile Pro Asn Val Ser Asp Glu Val Leu Ala Asn Leu Gly Glu Asp Gly Ile Ile Arg Ile Gly Ala Glu 805 810 Val Lys Pro Gly Asp Ile Leu Val Gly Lys Ile Thr Pro Lys Ser Glu 825 Thr Glu Leu Ala Pro Glu Glu Arg Leu Leu Arg Ala Ile Phe Gly Glu Lys Ala Ala Asp Val Lys Asp Ala Ser Leu Thr Val Pro Pro Gly Thr Glu Gly Val Val Met Asp Val Lys Val Phe Ser Arg Lys Asp Arg Leu Ser Lys Ser Asp Asp Glu Leu Val Glu Glu Ala Val His Leu Lys Asp Leu Gln Lys Gly Tyr Lys Asn Gln Val Ala Thr Leu Lys Thr Glu Tyr Arg Glu Lys Leu Gly Ala Leu Leu Leu Asn Glu Lys Ala Pro Ala Ala 920 Ile Ile His Arg Arg Thr Ala Glu Ile Val Val His Glu Gly Leu Leu Phe Asp Gln Glu Thr Ile Glu Arg Ile Glu Gln Glu Asp Leu Val Asp

945					950					955					960
Leu	Leu	Met	Pro	Asn 965	Cys	Glu	Met	Tyr	Glu 970	Val	Leu	Lys	Gly	Leu 975	Leu
Ser	Asp	Tyr	Glu 980	Thr	Ala	Leu	Gln	Arg 985	Leu	Glu	Ile	Asn	Tyr 990	Lys	Thr
Glu	Val	Glu 995	His	Ile	Arg	Glu	Gly 1000		Ala	Asp	Leu	Asp 1005		Gly	Val
Ile	Arg 1010		Val	Lys	Val	Tyr 1015		Ala	Ser	Lys	Arg 1020		Leu	Gln	Val
Gly 1025	Asp	Lys	Met	Ala	Gly 1030		His	Gly	Asn	Lys 1035		Val	Val	Ser	Lys 1040
Ile	Val	Pro	Glu	Ala 1045	-	Met	Pro	Tyr	Leu 1050		Asn	Gly	Glu	Thr 1055	
Gln	Met	Ile	Leu 1060		Pro	Leu	Gly	Val 1065		Ser	Arg	Met	Asn 1070		Gly
Gln	Val	Leu 1075		Thr	His	Leu	Gly 1080		Ala	Ala	Lys	Thr 1085		Gly	Ile
Tyr	Val 1090	-	Thr	Pro	Val	Phe 1095		Gly	Phe	Pro	Glu 1100		Arg	Ile	Trp
Asp 1105	Met	Met	Ile	Glu	Gln 1110		Leu	Pro	Glu -	Asp 1115	_	Lys	Ser	Phe	Leu 1120
Tyr	Asp	Gly	Lys	Thr 1125		Glu	Arg	Phe	Asp 1130		Lys	Val	Val	Ile 1135	-
Tyr	Ile	Tyr	Met 1140		Lys	Leu	Ser	His 1145		Ile	Ala	Asp	Lys 1150		His
Ala	Arg	Ser 1155		Gly	Pro	Tyr	Ser 1160		Val	Thr	Gln	Gln 1165		Leu	Gly
Gly	Lys 1170		Gln	Met	Gly	Gly 1175		Arg	Phe	Gly	Glu 1180		Glu	Val	Trp
Ala 1185	Leu	Glu	Ala	Tyr	Gly 1190			His	Met	Leu 1195		Glu	Ile	Leu	Thr 1200
Val	Lys	Ser	Asp	Asp 1205		Ser	Gly	Arg	Thr 1210	_	Ile	Tyr	Glu	Ser 1215	
Val	Lys	Gly	Glu 1220		Leu	Leu	Arg	Ser 1225		Thr	Pro	Glu	Ser 1230		Asn
Val	Leu	Ile 1235		Glu	Met	Gln	Gly 1240		Gly	Leu	Asp	Val 1245		Pro	Met
Val	Val 1250		Ala												

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<211> 224
<212> PRT
<213> Chlamydia pneumoniae
<400> 397
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Met Leu Lys His Glu Phe Tyr Gln Arg Trp Ser Glu Gly Lys Leu Glu
Lys Gln Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr Leu His Ile Lys
Ala Phe Pro Cys Tyr Leu Ser Ala Leu His Ala Arg Cys Asp Asp Leu
Gln Ile Arg Arg Gln Ile Leu Glu Asn Leu Met Asp Glu Glu Ala Gly
Asn Pro Asn His Ile Asp Leu Trp Arg Gln Phe Ala Leu Ser Leu Gly
Val Ser Glu Glu Leu Ala Asn His Glu Phe Ser Gln Ala Ala Gln
Asp Met Val Ala Thr Phe Arg Arg Leu Cys Asp Met Pro Gln Leu Ala
Val Gly Leu Gly Ala Leu Tyr Thr Tyr Glu Ile Gln Ile Pro Gln Val
Cys Val Glu Lys Ile Arg Gly Leu Lys Glu Tyr Phe Gly Val Ser Ala
Arg Gly Tyr Ala Tyr Phe Thr Val His Gln Glu Ala Asp Ile Lys His
Ala Ser Glu Glu Lys Glu Met Leu Gln Thr Leu Val Gly Arg Glu Asn
                                185
Pro Asp Ala Val Leu Gln Gly Ser Gln Glu Val Leu Asp Thr Leu Trp
Asn Phe Leu Ser Ser Phe Ile Asn Ser Thr Glu Pro Cys Ser Cys Lys
                        215
<210> 398
<211> 556
<212> PRT
<213> Chlamydia pneumoniae
<400> 398
Met Ser Lys Leu Ile Arg Arg Val Val Thr Val Leu Ala Leu Thr Ser
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Met Ala Ser Cys Phe Ala Ser Gly Gly Ile Glu Ala Ala Val Ala Glu 20 25 30

Ser Leu Ile Thr Lys Ile Val Ala Ser Ala Glu Thr Lys Pro Ala Pro

40 45 35 Val Pro Met Thr Ala Lys Lys Val Arg Leu Val Arg Arg Asn Lys Gln Pro Val Glu Gln Lys Ser Arg Gly Ala Phe Cys Asp Lys Glu Phe Tyr Pro Cys Glu Glu Gly Arg Cys Gln Pro Val Glu Ala Gln Gln Glu Ser Cys Tyr Gly Arg Leu Tyr Ser Val Lys Val Asn Asp Asp Cys Asn Val 105 Glu Ile Cys Gln Ser Val Pro Glu Tyr Ala Thr Val Gly Ser Pro Tyr Pro Ile Glu Ile Leu Ala Ile Gly Lys Lys Asp Cys Val Asp Val Val Ile Thr Gln Gln Leu Pro Cys Glu Ala Glu Phe Val, Ser Ser Asp Pro Glu Thr Thr Pro Thr Ser Asp Gly Lys Leu Val Trp Lys Ile Asp Arg Leu Gly Ala Gly Asp Lys Cys Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly Cys Cys Phe Thr Ala Ala Thr Val Cys Ala Cys Pro Glu Leu Arg Ser Tyr Thr Lys Cys Gly Gln Pro Ala Ile Cys Ile Lys Gln Glu Gly Pro Asp Cys Ala Cys Leu Arg Cys Pro Val Cys Tyr Lys Ile Glu Val Val Asn Thr Gly Ser Ala Ile Ala Arg Asn Val Thr Val Asp 250 Asn Pro Val Pro Asp Gly Tyr Ser His Ala Ser Gly Gln Arg Val Leu Ser Phe Asn Leu Gly Asp Met Arg Pro Gly Asp Lys Lys Val Phe Thr Val Glu Phe Cys Pro Gln Arg Arg Gly Gln Ile Thr Asn Val Ala Thr Val Thr Tyr Cys Gly Gly His Lys Cys Ser Ala Asn Val Thr Thr Val Val Asn Glu Pro Cys Val Gln Val Asn Ile Ser Gly Ala Asp Trp Ser Tyr Val Cys Lys Pro Val Glu Tyr Ser Ile Ser Val Ser Asn Pro Gly Asp Leu Val Leu His Asp Val Val Ile Gln Asp Thr Leu Pro Ser Gly 360

203

Val Thr Val Leu Glu Ala Pro Gly Gly Glu Ile Cys Cys Asn Lys Val 370 375 380

Val Trp Arg Ile Lys Glu Met Cys Pro Gly Glu Thr Leu Gln Phe Lys 385 390 395 400

Leu Val Val Lys Ala Gln Val Pro Gly Arg Phe Thr Asn Gln Val Ala
405 410 415

Val Thr Ser Glu Ser Asn Cys Gly Thr Cys Thr Ser Cys Ala Glu Thr 420 425 430

Thr Thr His Trp Lys Gly Leu Ala Ala Thr His Met Cys Val Leu Asp 435 440 445

Thr Asn Asp Pro Ile Cys Val Gly Glu Asn Thr Val Tyr Arg Ile Cys 450 460

Val Thr Asn Arg Gly Ser Ala Glu Asp Thr Asn Val Ser Leu Ile Leu 465 470 475 480

Lys Phe Ser Lys Glu Leu Gln Pro Ile Ala Ser Ser Gly Pro Thr Lys 485 490 495

Gly Thr Ile Ser Gly Asn Thr Val Val Phe Asp Ala Leu Pro Lys Leu 500 505 510

Gly Ser Lys Glu Ser Val Glu Phe Ser Val Thr Leu Lys Gly Ile Ala 515 520 525

Pro Gly Asp Ala Arg Gly Glu Ala Ile Leu Ser Ser Asp Thr Leu Thr 530 540

Ser Pro Val Ser Asp Thr Glu Asn Thr His Val Tyr 545 550 555

<210> 399

<211> 461

<212> PRT

<213> Chlamydia pneumoniae ·

<400> 399

Met Thr Gln Glu Phe Asp Cys Val Val Ile Gly Ala Gly Pro Ser Gly 5 10 15

Tyr Val Ala Ala Ile Thr Ala Ala Gln Ser Lys Leu Arg Thr Ala Leu 20 25 30

Ile Glu Glu Asp Gln Ala Gly Gly Thr Cys Leu Asn Arg Gly Cys Ile

Pro Ser Lys Ala Leu Ile Ala Gly Ala Asn Val Val Ser His Ile Lys 50 55 60

His Ala Glu Gln Phe Gly Ile His Val Asp Gly Tyr Thr Ile Asp Tyr 65 70 75 80

Pro Ala Met Ala Lys Arg Lys Asn Thr Val Val Gln Gly Ile Arg Gln

Gly	Leu	Glu	Gly 100	Leu	Ile	Arg	Ser	Asn 105	Lys	Ile	Thr	Val	Leu 110	Lys	Gly
Thr	Gly	Ser 115	Leu	Val	Ser	Ser	Thr 120	Glu	Val	Lys	Val	Ile 125	Gly	Gln	Asp
Thr	Thr 130	Ile	Ile	Lys	Ala	Asn 135	His	Ile	Ile	Leu	Ala 140	Thr	Gly	Ser	Glu
Pro 145	Arg	Pro	Phe	Pro	Gly 150	Val	Pro	Phe	Ser	Ser 155	Arg	Ile	Leu	Ser	Ser 160
Thr	Gly	Ile	Leu	Glu 165	Leu	Glu	Val	Leu	Pro 170	Lys	Lys	Leu	Ala	Ile 175	Ile
Gly	Gly	Gly	Val 180	Ile	Gly	Суз	Glu	Phe 185	Ala	Ser	Leu	Phe	His 190	Thr	Leu
Gly	Val	Glu 195	Ile	Thr	Val	Ile	Glu 200	Ala	Leu	Asp	His	Ile 205	Leu	Ala	Val
Asn	Asn 210	Lys	Glu	Val	Ser	Gln 215	Thr	Val	Thr	Asn	Lys 220	Phe	Thr	Lys	Gln
Gly 225	Ile	Arg	Ile	Leu	Thr 230	Lys	Ala	Ser	İle	Ser 235	Ala	Ile	Glu	Glu	Ser 240
Gln	Asn	Gln	Val	Arg 245	Ile	Thr	Val	Asn	Asp 250	Gln	Val	Glu	Glu	Phe 255	Asp
Tyr	Val	Leu	Val 260	Ala	Ile	Gly	Arg	Gln 265	Phe ·	Asn	Thr	Ala	Ser 270	Ile	Gly
Leu	Asp	Asn 275	Ala	Gly	Val	Ile	Arg 280	Asp	Asp	Arg	Gly	Val 285	Ile	Pro	Val
Asp	Glu 290	Thr	Met	Arg	Thr	Asn 295	Val	Pro	Asn	Ile	Tyr 300	Ala	Ile	Gly	Asp
Ile 305	Thr	Gly	Lys	Trp	Leu 310	Leu	Ala	His	Val	Ala 315	Ser	His	Gln	Gly	Val 320
Ile	Ala	Ala	Lys	Asn 325	Ile	Ser	Gly	His	His 330	Glu	Val	Met	Asp	Tyr 335	Ser
Ala	Ile	Pro	Ser 340	Val	Ile	Phe	Thr	His 345	Pro	Glu	Ile	Ala	Met 350	Val	Gly
Leu	Ser	Leu 355	Gln	Glu	Ala	Glu	Gln 360	Gln	Asn	Leu	Pro	Ala 365	Lys	Leu	Thr
Lys	Phe 370	Pro	Phe	Lys	Ala	Ile 375	Gly	Lys	Ala	Val	Ala 380	Leu	Gly	Ala	Ser
Asp 385	Gly	Phe	Ala	Ala	Ile 390	Val	Ser	His	Glu	Ile 395	Thr	Gln	Gln	Ile	Leu 400
Gly	Ala	Tyr	Val	Ile 405	Gly	Pro	His	Ala	Ser 410	Ser	Leu	Ile	Gly	Glu 415	Met

Thr Leu Ala Ile Arg Asn Glu Leu Thr Leu Pro Cys Ile Tyr Glu Thr 420 425 430

Val His Ala His Pro Thr Leu Ser Glu Val Trp Ala Glu Gly Ala Leu 435 440 445

Leu Ala Thr Asn His Pro Leu His Phe Pro Pro Lys Ser 450 455 460

<210> 400

<211> 544

<212> PRT

<213> Chlamydia pneumoniae

<400> 400

Met Ala Ala Lys Asn Ile Lys Tyr Asn Glu Glu Ala Arg Lys Lys Ile $5 \hspace{1.5cm} 10 \hspace{1.5cm} 15$

His Lys Gly Val Lys Thr Leu Ala Glu Ala Val Lys Val Thr Leu Gly 20 25 30

Pro Lys Gly Arg His Val Val Ile Asp Lys Ser Phe Gly Ser Pro Gln 35 40 45

Val Thr Lys Asp Gly Val Thr Val Ala Lys Glu Ile Glu Leu Glu Asp 50 60

Lys His Glu Asn Met Gly Ala Gln Met Val Lys Glu Val Ala Ser Lys 65 70 75 80

Thr Ala Asp Lys Ala Gly Asp Gly Thr Thr Thr Ala Thr Val Leu Ala 85 90 95

Glu Ala Ile Tyr Ser Glu Gly Leu Arg Asn Val Thr Ala Gly Ala Asn 100 105 110

Pro Met Asp Leu Lys Arg Gly Ile Asp Lys Ala Val Lys Val Val Val 115 120 125

Asp Glu Leu Lys Lys Ile Ser Lys Pro Val Gln His His Lys Glu Ile 130 135 140

Ala Gln Val Ala Thr Ile Ser Ala Asn Asn Asp Ser Glu Ile Gly Asn 145 150 155 160

Leu Ile Ala Glu Ala Met Glu Lys Val Gly Lys Asn Gly Ser Ile Thr 165 170 175

Val Glu Glu Ala Lys Gly Phe Glu Thr Val Leu Asp Val Val Glu Gly 180 185 190

Met Asn Phe Asn Arg Gly Tyr Leu Ser Ser Tyr Phe Ser Thr Asn Pro 195 200 205

Glu Thr Gln Glu Cys Val Leu Glu Asp Ala Leu Ile Leu Ile Tyr Asp 210 215 . 220

Lys Lys Ile Ser Gly Ile Lys Asp Phe Leu Pro Val Leu Gln Gln Val 225 230 235 240

PCT/US01/23121 WO 02/08267

Ala	Glu	Ser	Gly	Arg 245	Pro	Leu	Leu	Ile	Ile 250	Ala	Glu	Glu	Ile	Glu 255	Gly
Glu	Ala	Leu	Ala 260	Thr	Leu	Val	Val	Asn 265	Arg	Leu	Arg	Ala	Gly 270	Phe	Arg
Val	Cys	Ala 275	Val	Lys	Ala	Pro	Gly 280	Phe	Gly	Asp	Arg	Arg 285	Lys	Ala	Met
Leu	Glu 290	Asp	Ile	Ala	Ile	Leu 295	Thr	Gly	Gly	Gln	Leu 300	Val	Ser	Glu	Glu
Leu 305	Gly	Met	Lys	Leu	Glu 310	Asn	Thr	Thr	Leu	Ala 315	Met	Leu	Gly	Lys	Ala 320
Lys	Lys	Val	Ile	Val 325	Thr	Lys	Glu	Asp	Thr 330	Thr	Ile	Val	Glu	Gly 335	Leu
Gly	Asn	Lys	Pro 340	Asp	Ile	Gln	Ala	Arg 345	Cys	Asp	Asn	Ile	Lys 350	Lys	Gln
Ile	Glu	Asp 355	Ser	Thr	Ser	Asp	Туr 360	Asp	Lys	Glu	Lys	Leu 365	Gln	Glu	Arg
Leu	Ala 370	Lys	Leu	Ser	Gly	Gly 375	Val	Ala	Val	Ile	Arg 380	Val	Gly	Ala	Ala
Thr 385	Glu	Ile	Glu	Met	Lys 390	Glu	Lys	Lys	Asp	Arg 395		Asp	Asp	Ala	Gln 400
His	Ala	Thr	Ile	Ala 405	Ala	Val	Glu	Glu	Gly 410	Ile	Leu	Pro	Gly	Gly 415	Gly
Thr	Ala	Leu	Val 420	Arg	Cys	Ile	Pro	Thr 425	Leu	Glu	Ala	Phe	Leu 430	Pro	Met
Leu	Ala	Asn 435	Glu	Asp	Glu	Ala	Ile 440	Gly	Thr	Arg	Ile	11e 445	Leu	Lys	Ala
Leu	Thr 450	Ala	Pro	Leu	Lys	Gln 455	Ile	Ala	Ser	Asn	Ala 460	Gly	Lys	Glu	Gly
Ala 465	Ile	Ile	Cys	Gln	Gln 470	Val	Leu	Ala	Arg	Ser 475	Ala	Asn	Glu	Gly	Tyr 480
Asp	Ala	Leu	Arg	Asp 485	Ala	Tyr	Thr	Asp	Met 490	Ile	Asp	Ala	Gly	11e 495	Leu
Asp	Pro	Thr	Lys 500	Val	Thr	Arg	Ser	Ala 505	Leu	Glu	Ser	Ala	Ala 510	Ser	Ile
Ala	Gly	Leu 515	Leu	Leu	Thr	Thr	Glu 520	Ala	Leu	Ile	Ala	Asp 525	Ile	Pro	Glu
Glu	Lys 530	Ser	Ser	Ser	Ala	Pro 535	Ala	Met	Pro	Ser	Ala 540	Gly	Met	Asp	Tyr

<210> 401 <211> 664 <212> PRT

<213> Chlamydia pneumoniae

<400> 401

Met Glu Lys Val Ser Ser Tyr Pro Ser Val Pro Leu Pro Leu Gly Ala
5 10 15

Ser Lys Ile Ser Pro Asn Arg Tyr Arg Phe Ala Leu Tyr Ala Ser Gln 20 25 30

Ala Thr Glu Val Ile Leu Ala Leu Thr Asp Glu Asn Ser Glu Val Ile 35 40 45

Glu Val Pro Leu Tyr Pro Asp Thr His Arg Thr Gly Ala Ile Trp His 50 60

Ile Glu Ile Glu Gly Ile Ser Asp Gln Ser Ser Tyr Ala Phe Arg Val 65 70 75 80

His Gly Pro Lys Lys His Gly Met Gln Tyr Ser Phe Lys Glu Tyr Leu $85 \hspace{1.5cm} 90 \hspace{1.5cm} 95$

Ala Asp Pro Tyr Ala Lys Asn Ile His Ser Pro Gln Ser Phe Gly Ser 100 105 110

Arg Lys Gln Gly Asp Tyr Ala Phe Cys Tyr Leu Lys Glu Glu Pro 115 120 125

Phe Pro Trp Asp Gly Asp Gln Pro Leu His Leu Pro Lys Glu Glu Met 130 $$135\$

Ile Ile Tyr Glu Met His Val Arg Ser Phe Thr Gln Ser Ser Ser Ser 145 150 155 160

Arg Val His Ala Pro Gly Thr Phe Leu Gly Ile Ile Glu Lys Ile Asp 165 170 175

His Leu His Lys Leu Gly Ile Asn Ala Val Glu Leu Leu Pro Ile Phe 180 185 190

Glu Phe Asp Glu Thr Ala His Pro Phe Arg Asn Ser Lys Phe Pro Tyr 195 200 205

Leu Cys Asn Tyr Trp Gly Tyr Ala Pro Leu Asn Phe Phe Ser Pro Cys 210 215 220

Arg Arg Tyr Ala Tyr Ala Ser Asp Pro Cys Ala Pro Ser Arg Glu Phe 225 230 235 240

Lys Thr Leu Val Lys Thr Leu His Gln Glu Gly Ile Glu Val Ile Leu 245 250 255

Asp Val Val Phe Asn His Thr Gly Leu Gln Gly Thr Thr Cys Ser Leu 260 265 270

Pro Trp Ile Asp Thr Pro Ser Tyr Tyr Ile Leu Asp Ala Gln Gly His 275 280 285

Phe Thr Asn Tyr Ser Gly Cys Gly Asn Thr Leu Asn Thr Asn Arg Ala 290 295 300

Pro Thr Thr Gln Trp Ile Leu Asp Ile Leu Arg Tyr Trp Val Glu Glu

305					310					315					320
Met	His	Val	Asp	Gly 325	Phe	Arg	Phe	Asp	Leu 330	Ala	Ser	Val	Phe	Ser 335	Arg
Gly	Pro	Ser	Gly 340	Ser	Pro	Leu	Gln	Phe 345	Ala	Pro	Val	Leu	Glu 350	Ala	Ile
Ser	Phe	Asp 355	Pro	Leu	Leu	Ala	Ser 360	Thr	Lys	Ile	Ile	Ala 365	Glu	Pro	Trp
Asp	Ala 370	Gly	Gly	Leu	Tyr	Gln 375	Val	Gly	Tyr	Phe	Pro 380	Thr	Leu	Ser	Pro
Arg 385	Trp	Ser	Glu	Trp	Asn 390	Gly	Pro	Tyr	Arg	Asp 395	Asn	Val	Lys	Ala	Phe 400
Leu	Asn	Gly	Asp	Gln 405	Asn	Leu	Ile	Gly	Thr 410	Phe	Ala	Ser	Arg	Ile 415	Ser
Gly	Ser	Gln	Asp 420	Ile	Tyr	Pro	His	Gly 425	Ser	Pro	Thr	Asn	Ser 430	Ile	Asn
Tyr	Val	Ser 435	Cys	His	Asp	Gly	Phe 440	Thr	Leu	Cys	Asp	Thr 445	Val	Thr	Tyr
Asn	His 450	Lys	His	Asn	Glu	Ala 455	Asn	Gly	Glu	Asp	Asn 460	Arg	Asp	Gly	Thr
Asp 465	Ala	Asn	Tyr	Ser	Tyr 470	Asn	Phe	Gly	Thr	Glu 475	Gly	Lys	Thr	Glu	Asp 480
Pro	Gly	Ile	Leu	Glu 485	Val	Arg	Glu	Arg	Gln 490	Leu	Arg	Asn	Phe	Phe 495	Leu
Thr	Leu	Met	Val 500	Ser	Gln	Gly	Ile	Pro 505	Met	Ile	Gln	Ser	Gly 510	Asp	Glu
Tyr	Ala	His 515	Thr	Ala	Glu	Gly	Asn 520	Asn	Asn	Arg	Trp	Ala 525	Leu	Asp	Ser
Asn	Ala 530	Asn	Tyr	Phe	Leu	Trp 535	Asp	Gln	Leu	Thr	Ala 540	Lys	Pro	Thr	Leu
Met 545	His	Phe	Leu	Cys	Asp 550	Leu	Ile	Ala	Phe	Arg 555	Lys	Lys	Tyr	Lys	Thr 560
Leu	Phe	Asn	Arg	Gly 565	Phe	Leu	Ser	Asn	Lys 570	Glu	Ile	Ser	Trp	Val 575	Asp
Ala	Met	Gly	Asn 580	Pro	Met	Thr	Trp	Arg 585	Pro	Gly	Asn	Phe	Leu 590	Ala	Phe
Lys	Ile	Lys 595		Pro	Lys	Ala	His 600	Val	Tyr	Val	Ala	Phe 605	His	Val	Gly
Ala	Gln 610	Asp	Gln	Leu	Ala	Thr 615	Leu	Pro	Lys	Ala	Ser 620	Ser	Asn	Phe	Leu
Pro 625	Tyr	Gln	Ile	Val	Ala 630	Glu	Ser	Gln	Gln	Gly 635	Phe	Val	Pro	Gln	Asn 640

Val Ala Thr Pro Thr Val Ser Leu Gln Pro His Thr Thr Leu Ile Ala 645 650 655

Ile Ser His Ala Lys Glu Val Thr 660

<210> 402

<211> 328

<212> PRT

<213> Chlamydia pneumoniae

<400> 402

Met Ala Phe Lys Glu Val Val Arg Val Ala Val Thr Gly Gly Lys Gly
5 10 15

Gln Ile Ala Tyr Asn Phe Leu Phe Ala Leu Ala His Gly Asp Val Phe 20 25 30

Gly Val Asp Arg Gly Val Asp Leu Arg Ile Tyr Asp Val Pro Gly Thr 35 40 45

Glu Arg Ala Leu Ser Gly Val Arg Met Glu Leu Asp Asp Gly Ala Tyr 50 60

Pro Leu Leu His Arg Leu Arg Val Thr Thr Ser Leu Asn Asp Ala Phe 65 70 75 80

Asp Gly Ile Asp Ala Ala Phe Leu Ile Gly Ala Val Pro Arg Gly Pro 85 90 95

Gly Met Glu Arg Gly Asp Leu Leu Lys Gln Asn Gly Gln Ile Phe Ser 100 105 110

Leu Gln Gly Ala Ala Leu Asn Thr Ala Ala Lys Arg Asp Ala Lys Ile 115 120 125

Phe Val Val Gly Asn Pro Val Asn Thr Asn Cys Trp Ile Ala Met Lys 130 135 140

His Ala Pro Arg Leu His Arg Lys Asn Phe His Ala Met Leu Arg Leu 145 150 155 160

Asp Gln Asn Arg Met His Ser Met Leu Ala His Arg Ala Glu Val Pro 165 170 175

Leu Glu Glu Val Ser Arg Val Val Ile Trp Gly Asn His Ser Ala Lys 180 185 190

Gln Val Pro Asp Phe Thr Gln Ala Arg Ile Ser Gly Lys Pro Ala Ala 195 200 205

Glu Val Ile Gly Asp Arg Asp Trp Leu Glu Asn Ile Leu Val His Ser 210 215 220

Val Gln Asn Arg Gly Ser Ala Val Ile Glu Ala Arg Gly Lys Ser Ser 225 230 235 240

Ala Ala Ser Ala Ser Arg Ala Leu Ala Glu Ala Ala Arg Ser Ile Phe 245 250 255

Cys Pro Lys Ser Asp Glu Trp Phe Ser Ser Gly Val Cys Ser Asp His 260 265 270

Asn Pro Tyr Gly Ile Pro Glu Asp Leu Ile Phe Gly Phe Pro Cys Arg 275 280 285

Met Leu Pro Ser Gly Asp Tyr Glu Ile Ile Pro Gly Leu Pro Trp Glu 290 295 300

Pro Phe Ile Arg Asn Lys Ile Gln Ile Ser Leu Asp Glu Ile Ala Gln 305 310 315 320

Glu Lys Ala Ser Val Ser Ser Leu 325

<210> 403

<211> 217

<212> PRT

<213> Chlamydia pneumoniae

<400> 403

Met Lys Arg Val Ile Tyr Lys Thr Ile Phe Cys Gly Leu Thr Leu Leu 5 10

Thr Ser Leu Ser Ser Cys Ser Leu Asp Pro Lys Gly Tyr Asn Leu Glu 20 25 30

Thr Lys Asn Ser Arg Asp Leu Asn Gln Glu Ser Val Ile Leu Lys Glu 35 40 45

Asn Arg Glu Thr Pro Ser Leu Val Lys Arg Leu Ser Arg Arg Ser Arg 50 55 60

Arg Leu Phe Ala Arg Arg Asp Gln Thr Gln Lys Asp Thr Leu Gln Val 65 70 75 80

Gln Ala Asn Phe Lys Thr Tyr Ala Glu Lys Ile Ser Glu Gln Asp Glu 85 90 95

Arg Asp Leu Ser Phe Val Val Ser Ser Ala Ala Glu Lys Ser Ser Ile 100 105 110

Ser Leu Ala Leu Ser Gln Gly Glu Ile Lys Asp Ala Leu Tyr Arg Ile 115 120 125

Arg Glu Val His Pro Leu Ala Leu Ile Glu Ala Leu Ala Glu Asn Pro 130 135 140

Ala Leu Ile Glu Gly Met Lys Lys Met Gln Gly Arg Asp Trp Ile Trp 145 150 155 160

Asn Leu Phe Leu Thr Gln Leu Ser Glu Val Phe Ser Gln Ala Trp Ser 165 170 175

Gln Gly Val Ile Ser Glu Glu Asp Ile Ala Ala Phe Ala Ser Thr Leu 180 185 190

Gly Leu Asp Ser Gly Thr Val Ala Ser Ile Val Gln Gly Glu Arg Trp
195 200 205

Pro Glu Leu Val Asp Ile Val Ile Thr 210 215

<210> 404

<211> 270

<212> PRT

<213> Chlamydia pneumoniae

<400> 404

Met Ile Ile Lys Asn Asn Glu Leu Met Ile Arg Arg Phe Phe Lys
5 10 15

Thr Leu Phe Pro Pro Gly Pro Gln Tyr Ser Leu Cys Tyr Ala Ser Ile 20 25 30

Leu Ile Val Leu Ser Ser Leu Val Cys Val Pro Thr Phe Cys Trp Leu 35 40 45

Phe Leu Pro Glu Leu Ser Leu Ser Lys Phe Asn Pro Ser Pro Ile Arg 50 55 60

Asn Leu Phe Leu Val Ser Ser Thr Leu Ser Lys Val Pro Pro Thr Ala 65 70 75 80

Ile Ala Glu His Leu Arg Leu Ser Ala Asp Ala Pro Thr Tyr Leu His
85 90 95

Glu Phe Ser Iİe Lys Glu Ala Glu Ser Ser Leu His Ala Leu Gly Ile 100 105 110

Phe Ser Ser Leu Val Ile Glu Lys Ser Pro Asp Asn Lys Gly Ile Thr

Ile Phe Tyr Thr Leu Gln Thr Pro Ile Ala Tyr Val Gly Asn Arg Ser 130 135 140

Asn Thr Leu Cys Asn Leu Glu Gly Ser Cys Phe Leu Gly Gln Pro Tyr 145 150 155 160

Phe Pro Ser Leu Asn Leu Pro Gln Ile Phe Phe Ser Gln Glu Asp Leu 165 170 175

Lys Met Gln Lys Leu Pro Lys Glu Lys Met Leu Phe Thr Lys Ile Leu 180 185 190

Leu Lys Glu Leu Ala Met Glu Ser Pro Lys Ile Ile Asp Leu Ser Leu 195 200 205

Ser Asp Ala Tyr Pro Gly Glu Ile Ile Val Thr Leu Ser Ser Gly Ser 210 220

Leu Leu Arg Leu Pro Ile Lys Thr Leu Asp Arg Ala Leu Asp Leu Tyr 225 230 235 240

Lys His Met Lys Lys Ser Pro Val Ile Glu Ser Glu Lys Gln Tyr Val 245 250 255

Tyr Asp Leu Arg Phe Pro Asn Phe Leu Leu Leu Lys Ala Leu 260 265 270

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214

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WO 02/08267

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325 330 335 Asp Met Glu Asp Phe Asp Pro Ser Gly Pro Pro Pro Trp Glu Glu Phe 345 Ala Lys Ile Ile Gln Val Phe Ser Ser Asn Thr Glu Ala Leu Ile Ile Asp Gln Thr Asn Asn Pro Gly Gly Ser Val Leu Tyr Leu Tyr Ala Leu Leu Ser Met Leu Thr Asp Arg Pro Leu Glu Leu Pro Lys His Arg Met Ile Leu Thr Gln Asp Glu Val Val Asp Ala Leu Asp Trp Leu Thr Leu Leu Glu Asn Val Asp Thr Asn Val Glu Ser Arg Leu Ala Leu Gly Asp Asn Met Glu Gly Tyr Thr Val Asp Leu Gln Val Ala Glu Tyr Leu Lys Ser Phe Gly Arg Gln Val Leu Asn Cys Trp Ser Lys Gly Asp Ile Glu Leu Ser Thr Pro Ile Pro Leu Phe Gly Phe Glu Lys Ile His Pro His Pro Arg Val Gln Tyr Ser Lys Pro Ile Cys Val Leu Ile Asn Glu Gln Asp Phe Ser Cys Ala Asp Phe Phe Pro Val Val Leu Lys Asp Asn Asp Arg Ala Leu Ile Val Gly Thr Arg Thr Ala Gly Ala Gly Gly Phe Val 520 Phe Asn Val Gln Phe Pro Asn Arg Thr Gly Ile Lys Thr Cys Ser Leu Thr Gly Ser Leu Ala Val Arg Glu His Gly Ala Phe Ile Glu Asn Ile Gly Val Glu Pro His Ile Asp Leu Pro Phe Thr Ala Asn Asp Ile Arg Tyr Lys Gly Tyr Ser Glu Tyr Leu Asp Lys Val Lys Lys Leu Val Cys

Gln Leu Ile Asn Asn Asp Gly Thr Ile Ile Leu Ala Glu Asp Gly Ser 595 600 605

Phe

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<213> Chlamydia trachomatis serovar D

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15

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WO 02/08267 PCT/US01/23121

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His Pro His Gly Asp Ala Pro Ile Val Glu Ala Leu Val Val Leu Ala

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Thr	Gly	Asp	Pro 100	His	Ala	Ala	Ala	Arg 105	Tyr	Ile	Glu	Ala	Arg 110	Leu	Ser
Pro		Ala 115	Lys	Glu	Val	Leu	Phe 120	Asn	Thr	Asp	Leu	Met 125	Thr	Phe	His
Asp	Ser 130	Tyr	Asp	Gly	Arg	Glu 135	Gln	Glu	Pro	Asp	Ile 140	Leu	Ala	Ala	Lys
Ile 145	Pro	Leu	Leu	Ļeu	Leu 150	His	Gly	Val	Asp	Gly 155	Ile	Ala	Val	Gly	Met 160
Thr	Thr	Lys	Ile	Phe 165	Pro	His	Asn	Phe	Cys 170	Asp	Leu	Leu	Glu	Ala 175	Gln
Ile	Ala	Ile	Leu [.] 180	Asn	Asp	Gln	Pro	Phe 185	Ser	Leu	Leu	Pro	Asp 190	Phe	Pro
Pro	Gly	Gly 195	Thr	Met	Asp	Ala	Ser 200	Asp	Tyr	Gln	Asp	Gly 205	Leu	Gly	Ser
Ile	Val 210	Leu	Arg	Ala	Thr	Ile 215	Asp	Ile	Ile	Asn	Asp 220	Lys	Thr	Leu	Leu
Ile 225	Lys	Glu	Ile	Cys	Pro 230	Ser	Thr	Thr	Thr	Glu 235	Thr	Leu	Ile	Arg	Ser 240
Ile	Glu	Asn	Ala	Ala 245	Lys	Arg	Gly	Ile	Ile 250	Lys	Ile	Asp	Ser	Ile 255	Gln
Asp	Phe	Ser	Thr 260	Asp	Leu	Pro	His	Ile 265	Glu	Ile	Lys	Leu	Pro 270	Lys	Gly
Ile	Tyr	Ala 275	Lys	Asp	Leu	Leu	Arg 280	Pro	Leu	Tyr	Thr	His 285	Thr	Glu	Cys
Gln	Val 290	Ile	Leu	Thr	Ser	Arg 295	Pro	Thr	Ala	Ile	Tyr 300	Gln	Gly	Lys	Pro
Trp 305	Glu	Thr	Thr	Ile	Ser 310	Glu	Ile	Leu	Arg	Leu 315	Gln	Thr	Lys	Thr	Leu 320
Gln	Asn	Tyr	Leu	Lуs 325	Lys	Glu	Leu	Leu	Ile 330	Leu	Glu	Asp	Ser	Leu 335	Ser
Arg	Glu	Leu	Tyr 340	His	Lys	Thr	Leu	Glu 345	Tyr	Leu	Phe	Ile	Lys 350	His	Lys
Leu	Tyr	Asp 355	Thr	Val	Arg	Ser	Met 360	Leu	Ser	Lys	Arg	Lys 365	Thr	Ser	Pro
Ser	Ser 370	Ser	Thr	Ile	His	Asn 375	Ala	Val	Leu	Glu	Ala 380	Leu	Thr	Pro	Phe
Leu 385	Asp	Thr	Leu	Pro	Ala 390	Pro	Asp	Lys	Gln	Ala 395	Thr	Ala	Gln	Leu	Ala 400

Ala Leu Thr Ile Lys Lys Ile Leu Cys Phe Asp Glu Asn Ser Tyr Glu
405 410 415

Lys Glu Leu Ala Cys Leu Glu Lys Lys Arg Ser Ser Val Gln Lys Asp

420 425 430

Leu Ser Gln Leu Lys Lys Tyr Thr Val Leu Tyr Ile Lys Lys Leu Leu 435 440 445

Glu Thr Tyr Arg Gln Leu Gly His Arg Lys Thr Lys Ile Ala Lys Phe 450 455 460

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Leu Ala Ala Leu Asp Gln Glu Glu Asn Phe 485 490

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<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 435

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Glu Glu Val His Leu Lys Glu Val Gln Gly Thr Asn Thr Ile Ile Tyr $20 \hspace{1.5cm} 25 \hspace{1.5cm} 30$

Glu Leu Thr Val Ala Lys Gly Asp Ile Gly Lys Ile Ile Gly Lys Glu
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<213> Chlamydia trachomatis serovar D

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Thr Lys Glu Ser Glu Ala Ser Pro Ser Ala Ser Ser Ser Val Ser Ser 35 40 45

Trp Ser Phe Leu Ser Ser Ala Lys His Ala Leu Ile Ser Leu Arg Asp 50 60

Ala Ile Leu Asn Lys Asn Ser Ser Pro Thr Asp Ser Leu Ser Gln Leu

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Tyr	Asn	Glu	Ala 100	Lys	Ser	Asn	Phe	Asp 105	Thr	Ala	Lys	Ser	Gly 110	Leu	Glu
Asn	Ala	Thr 115	Thr	Leu	Ala	Glu	Tyr 120	Glu	Thr	Lys	Met	Ala 125	Asp	Leu	Met
Ala	Ala 130	Leu	Gln	Asp	Met	Glu 135	Arg	Leu	Ala	Lys	Gln 140	Lys	Ala	Glu	Val
Thr 145	Arg	Ile	Lys	Glu	Ala 150	Leu	Gln	Glu	Lys	Gln 155	Glu	Val	Ile	Asp	Lys 160
Leu	Asn	Gln	Leu	Val 165	Lys	Leu	Glu	Lys	Gln 170	Asn	Gln	Thr	Leu	Lys 175	Glu
Thr	Leu	Thr	Thr 180	Thr	Asp	Ser	Ala	Asp 185	Gln	Ile	Pro	Ala	Ile 190	Asn	Ser
Gln	Leu	Glu 195	Ile	Asn	Lys	Asn	Ser 200	Ala	Asp	Gln	Ile	Ile 205	Lys	Asp	Leu
Glu	Gly 210	Gln	Asn	Ile	Ser	Tyr 215	Glu	Ala	Val	Leu	Thr 220	Asn	Ala	Gly	Glu
Val 225	Ile	Lys	Ala	Ser	Ser 230	Glu	Ala	Gly	Ile	Lys 235	Leu	Gly	Gln	Ala	Leu 240
Gln	Ser	Ile	Val	Asp 245	Ala	Gly	Asp	Gln	Ser 250	Gln	Ala	Ala	Val	Leu 255	Gln
Ala	Gln	Gln	Asn 260	Asn	Ser	Pro	Asp	Asn 265	Ile	Ala	Ala	Thr	Lys 270	Lys	Leu
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Gly	Leu 290	Thr	Asp	Ser	Pro	Leu 295	Val	Lys	Lys	Ala	Glu 300	Glu	Gln	Ile	Ser
Gln 305	Ala	Gln	Lys	Asp	Ile 310	Gln	Glu	Ile	Lys	Pro 315	Ser	Gly	Ser	Asp	Ile 320
Pro	Ile	Val	Gly	Pro 325	Ser	Gly	Ser	Ala	Ala 330	Ser	Ala	Gly	Ser	Ala 335	Val
Gly	Ala	Leu	Lys 340	Ser	Ser	Asn	Asn	Ser 345	Gly	Arg	Ile	Ser	Leu 350	Leu	Leu
Asp	Asp	Val 355	Asp	Asn	Glu	Met	Ala 360	Ala	Ile	Ala	Met	Gln 365	Gly	Phe	Arg
Ser	Met 370	Ile	Glu	Gln	Phe	Asn 375	Val	Asn	Asn	Pro	Ala 380	Thr	Ala	Lys	Glu
Leu 385	Gln	Ala	Met	Glu	Ala 390	Gln	Leu	Thr	Ala	Met 395	Ser	Asp	Gln	Leu	Val 400

Gly	Ala	Asp	Gly	Glu 405	Leu	Pro	Ala	Glu	Ile 410	Gln	Ala	Ile	Lys	Asp 415	Ala
Leu	Ala	Gln	Ala 420	Leu	Lys	Gln	Pro	Ser 425	Thr	Asp	Gly	Leu	Ala 430	Thr	Ala
Met	Gly	Gln 435	Val	Ala	Phe	Ala	Ala 440	Ala	Lys	Val	Gly	Gly 445	Gly	Ser	Ala
Gly	Thr 450	Ala	Gly	Thr	Val	Gln 455	Met	Asn	Val	Lys	Gln 460	Leu	Tyr	Lys	Thr
Ala 465	Phe	Ser	Ser	Thr	Ser 470	Ser	Ser	Ser	Tyr	Ala 475	Ala	Ala	Leu	Ser	Asp 480
Gly	Tyr	Ser	Ala	Tyr 485	Lys	Thr	Leu	Asn	Ser 490	Leu	Tyr	Ser	Glu	Ser 495	Arg
Ser	Gly	Val	Gln 500	Ser	Ala	Ile	Ser	Gln 505	Thr	Ala	Asn	Pro	Ala 510	Leu	Ser
Arg	Ser	Val 515	Ser	Arg	Ser	Gly	Ile 520	Glu	Ser	Gln	Gly	Arg 525	Ser	Ala	Asp
Ala	Ser 530	Gln	Arg	Ala	Ala	Glu 535	Thr	Ile	Val	Arg	Asp 540	Ser	Gln	Thr	Leu
Gly 545	Asp	Val	Tyr	Ser	Arg 550	Leu	Gln	Val	Leu	Asp 555	Ser	Leu	Met	Ser	Thr 560
Ile	Val	Ser	Asn	Pro 565	Gln	Val	Asn	Gln	Glu 570	Glu	Île	Met	Gln	Lys 575	Leu
Thr	Ala	Ser	Ile 580	Ser	Lys	Ala	Pro	Gln 585	Phe	Gly	Tyr	Pro	Ala 590	۷al	Gln
Asn	Ser	Ala 595	dsÞ	Ser	Leu	Gln	L ys 600	Phe	Ala	Ala	Gln	Leu 605	Glu	Arg	Glu
Phe	Val 610	Asp	Gly	Glu	Arg	Ser 615	Leu	Ala	Glu	Ser	Arg 620	Glu	Asn	Ala	Phe
Arg 625	Lys	Gln	Pro	Ala	Phe 630	Ile	Gln	Gln	Val	Leu 635	Val	Asn	Ile	Ala	Ser 640
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Gln Asn Lys His Met Leu Glu His Thr Phe Tyr Val Lys Trp Ser Lys
20 25 30

<400> 437

Gly Glu Leu Thr Lys Glu Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr Leu His Ile Lys Ala Phe Pro Lys Tyr Leu Ser Ala Ile His Ser Arg Cys Asp Asp Leu Glu Ala Arg Lys Leu Leu Leu Asp Asn Leu Met Asp Glu Glu Asn Gly Tyr Pro Asn His Ile Asp Leu Trp Lys Gln Phe Val Phe Ala Leu Gly Val Thr Pro Glu Glu Leu Glu Ala His Glu Pro Ser Glu Ala Ala Lys Ala Lys Val Ala Thr Phe Met Arg Trp Cys Thr Gly Asp Ser Leu Ala Ala Gly Val Ala Ala Leu Tyr Ser Tyr Glu Ser Gln 135 Ile Pro Arg Ile Ala Arg Glu Lys Ile Arg Gly Leu Thr Glu Tyr Phe Gly Phe Ser Asn Pro Glu Asp Tyr Ala Tyr Phe Thr Glu His Glu Glu Ala Asp Val Arg His Ala Arg Glu Glu Lys Ala Leu Ile Glu Met Leu Leu Lys Asp Asp Ala Asp Lys Val Leu Glu Ala Ser Gln Glu Val Thr Gln Ser Leu Tyr Gly Phe Leu Asp Ser Phe Leu Asp Pro Gly Thr Cys Cys Ser Cys His Gln Ser Tyr <210> 438 <211> 533 <212> PRT <213> Chlamydia trachomatis serovar D Met Ser Asn Ser Phe Arg Asp Gln Glu Gln Gly Leu Gln Ala Val Phe Arg Ala Ala Arg Val Ile Ser His Met Phe Ser Gln Thr Ile Gly Pro Tyr Gly Phe Ser Thr Ile Val His Asn Val Gln Asp Thr Arg Thr Thr Gln Asp Ser Gln Ser Met Leu Lys Asp Ile Leu Phe Pro Asp Val Phe Glu Asn Ile Gly Met Lys Leu Ile Arg Asp Thr Ala Leu Arg Thr Arg 237

Met Arg Phe Gly Asp Gly Ala Lys Thr Thr Ala Leu Leu Ile Glu Ala Leu Leu Ala Glu Gly Met Thr Gly Ile Gln Lys Gly Leu Asp Pro His Glu Ile His Arg Gly Met Leu Leu Ala Glu Lys Lys Ile Gln Glu Val Phe Tyr Arg Glu Thr Phe Pro Leu Ser Asp Leu Glu His Thr Val Tyr Val Ser Ser Ile Ala Arg Arg Cys Asn Ser Glu Ile Ala Ser Val Leu 150 Ser Ser Ala Val Gly Tyr Gly Gly Lys Asn Gly Tyr Tyr Ile Val Glu Glu His Glu Glu His Glu Thr Tyr Trp His Ala Glu Glu His Ala Val 185 Trp Asp Phe Gly Tyr Ala Ser Pro Tyr Phe Ile Thr His Ala Glu Thr Gly Thr Val Glu Tyr Ser Gln Val Tyr Ile Leu Val Ser Glu Gln Pro 215 . Leu His Tyr Ser Asn Pro Ser Phe Leu Thr Phe Leu Gln Ser Val Val Gln Ala Gly Lys Thr Pro Leu Val Ile Leu Ala Glu Ala Phe Asp Lys 250 Glu Leu Leu Ala Met Leu Glu Met Asn Gln Ile Glu Arg Val Phe Pro Val Cys Ala Val Lys Val Ser Gly Lys His Ala Arg Glu Ser Leu Glu Asp Ile Ala Val Leu Thr Gly Ala Thr Leu Leu Ser Glu Met Asp Phe Glu Asp Ser Glu Glu Glu Arg Ile Thr Asn Arg Leu Gly Phe Val Ala Gly Ile Cys Val Ser Ser Thr Ser Leu Cys Val Pro Arg Glu Thr Asp 330 Asn Lys Gln Arg Met Ala Glu His Cys Ala Phe Leu Gln Asp Lys Leu 345 Ser Phe Ser Gln Glu Glu Glu Ala Ser Ala Arg Leu Arg Arg Leu 360 Ala Arg Leu Ser Ser Gly Glu Val Cys Ile His Ile Ala Ala Asp Cys Ile Pro Gln Glu Glu Ile Gly Tyr Ile Thr Ser Ser Ile Arg Ala Met 390 395

Thr Glu Ser Leu Arg Ser Gly Cys Leu Pro Gly Gly Gly Cys Ala Phe Ile Arg Ala Ala Arg Glu Ile Ser Val Pro Leu Ala Leu Ser Pro Ser 425 Glu Arg Phe Gly Phe Leu Ala Val Leu Ser Ala Ala Glu Lys Pro Phe Arg Ala Ile Val Thr Arg Ser Arg Arg Val Glu Glu Val Phe Ser Glu Val Phe Ser Gln Ala Asp Trp Arg Val Gly Phe Asn Gly Val Ser Gly Phe Val Glu Asp Ile Val Ser Gln Gly Ile Cys Asp Gly Ala Ser Cys Ile Gln Tyr Ala Leu Ser His Ala Val Gly Thr Thr Gly Leu Leu Leu Thr Ser Ala Leu Phe Ile Ala Ser Gln Glu Pro Met Leu Arg Glu Glu Asn Ser Glu Glu 530 <210> 439 <211> 465 <212> PRT <213> Chlamydia trachomatis serovar D <400> 439 Met Asn Glu Ala Phe Asp Cys Val Val Ile Gly Ala Gly Pro Gly Gly Tyr Val Ala Ala Ile Thr Ala Ala Gln Ala Gly Leu Lys Thr Ala Leu Ile Glu Lys Arg Glu Ala Gly Gly Thr Cys Leu Asn Arg Gly Cys Ile Pro Ser Lys Ala Leu Leu Ala Gly Ala Glu Val Val Thr Gln Ile Arg His Ala Asp Gln Phe Gly Ile His Val Glu Gly Phe Ser Ile Asn Tyr 65 70 75 80 Pro Ala Met Val Gln Arg Lys Asp Ser Val Val Arg Ser Ile Arg Asp Gly Leu Asn Gly Leu Ile Arg Ser Asn Lys Ile Thr Val Phe Ser Gly Arg Gly Ser Leu Ile Ser Ser Thr Glu Val Lys Ile Leu Gly Glu Asn Pro Ser Val Ile Lys Ala His Ser Ile Ile Leu Ala Thr Gly Ser Glu 130

Pro Arg Ala Phe Pro Gly Ile Pro Phe Ser Ala Glu Ser Pro Arg Ile 150 Leu Cys Ser Thr Gly Val Leu Asn Leu Lys Glu Ile Pro Gln Lys Met 170 Ala Ile Ile Gly Gly Gly Val Ile Gly Cys Glu Phe Ala Ser Leu Phe His Thr Leu Gly Ser Glu Val Ser Val Ile Glu Ala Ser Ser Gln Ile Leu Ala Leu Asn Asn Pro Asp Ile Ser Lys Thr Met Phe Asp Lys Phe 215 Thr Arg Gln Gly Leu Arg Phe Val Leu Glu Ala Ser Val Ser Asn Ile Glu Asp Ile Gly Asp Arg Val Arg Leu Thr Ile Asn Gly Asn Val Glu 250 Glu Tyr Asp Tyr Val Leu Val Ser Ile Gly Arg Arg Leu Asn Thr Glu Asn Ile Gly Leu Asp Lys Ala Gly Val Ile Cys Asp Glu Arg Gly Val Ile Pro Thr Asp Ala Thr Met Arg Thr Asn Val Pro Asn Ile Tyr Ala Ile Gly Asp Ile Thr Gly Lys Trp Gln Leu Ala His Val Ala Ser His Gln Gly Ile Ile Ala Ala Arg Asn Ile Ala Gly His Lys Glu Glu Ile Asp Tyr Ser Ala Val Pro Ser Val Ile Phe Thr Phe Pro Glu Val Ala Ser Val Gly Leu Ser Pro Thr Ala Ala Gln Gln Lys Ile Pro Val 360 Lys Val Thr Lys Phe Pro Phe Arg Ala Ile Gly Lys Ala Val Ala Met Gly Glu Ala Asp Gly Phe Ala Ala Ile Ile Ser His Glu Thr Thr Gln 395 Gln Ile Leu Gly Ala Tyr Val Ile Gly Pro His Ala Ser Ser Leu Ile Ser Glu Ile Thr Leu Ala Val Arg Asn Glu Leu Thr Leu Pro Cys Ile Tyr Glu Thr Ile His Ala His Pro Thr Leu Ala Glu Val Trp Ala Glu Ser Ala Leu Leu Ala Val Asp Thr Pro Leu His Met Pro Pro Ala Lys 455

Lys

465

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WO 02/08267

<211> 122

<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 440

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240

Ile Ser Leu Thr Tyr Ile Tyr Gly Ile Gly Pro Ala Leu Ser Lys Glu 20 25 30

Ile Ile Ala Arg Leu Gln Leu Asn Pro Glu Ala Arg Ala Ala Glu Leu 35 40 45

Thr Glu Glu Glu Val Gly Arg Leu Asn Ala Leu Leu Gln Ser Asp Tyr 50 55 60

Val Val Glu Gly Asp Leu Arg Arg Val Gln Ser Asp Ile Lys Arg 65 70 75 80

Leu Ile Thr Ile His Ala Tyr Arg Gly Gln Arg His Arg Leu Ser Leu 85 90 95

Pro Val Arg Gly Gln Arg Thr Lys Thr Asn Ser Arg Thr Arg Lys Gly 100 105 110

Lys Arg Lys Thr Val Ala Gly Lys Lys 115 120

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<211> 553

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<213> Chlamydia trachomatis serovar D

<400> 441

Met Arg Ile Gly Asp Pro Met Asn Lys Leu Ile Arg Arg Ala Val Thr 5 10 15

Ile Phe Ala Val Thr Ser Val Ala Ser Leu Phe Ala Ser Gly Val Leu 20 25 30

Glu Thr Ser Met Ala Glu Ser Leu Ser Thr Asn Val Ile Ser Leu Ala 35 40 45

Asp Thr Lys Ala Lys Asp Asn Thr Ser His Lys Ser Lys Lys Ala Arg 50 60

Lys Asn His Ser Lys Glu Thr Pro Val Asp Arg Lys Glu Val Ala Pro 65 70 75 80

Val His Glu Ser Lys Ala Thr Gly Pro Lys Gln Asp Ser Cys Phe Gly

Arg Met Tyr Thr Val Lys Val Asn Asp Asp Arg Asn Val Glu Ile Thr 100 105 110

Gln	Ala	Val 115	Pro	Glu	Tyr	Ala	Thr 120	Val	Gly	Ser	Pro	Tyr 125	Pro	Ile	Glu
Iļe	Thr 130	Ala	Thr	Gly	Lys	Arg 135	Asp	Суѕ	Val	Asp	Val 140	Ile	Ile	Thr	Glņ
Gln 145	Leu	Pro	Cys	Glu	Ala 150	Glu	Phe	Val	Arg	Ser 155	Asp	Pro	Ala	Thr	Thr 160
Pro	Thr	Alá	Asp	Gly 165	Lys	Leu	Val	Trp	Lys 170	Ile	Asp	Arg	Leu	Gly 175	Gln
Gly	Glu	Lys	Ser 180	Lys	Ile	Thr	Val	Trp 185	Val	Lys	Pro	Leu	Lys 190	Glu	Gly
Cys	Суз	Phe 195	Thr	Ala	Ala	Thr	Val 200	Cys	Ala	Cys	Pro	Glu 205	Ile	Arg	Ser
Val	Thr 210	Lys	Суѕ	Gly	Gln	Pro 215	Ala	Ile	Cys	Val	Lys 220	Gln	Glu	Gly	Pro
Glu 225	Asn	Ala	Cys	Leu	Arg 230	Cys	Pro	Val	Val	Tyr 235	Lys	Ile	Asn	Ile	Val 240
Asn	Gln	Gly	Thr	Ala 245	Thr	Ala	Arg	Asn	Val 250	Val	Val	Glu	Asn	Pro 255	Val
Pro	Asp	Gly	Tyr 260	Ala	His	Ser	Ser	Gly 265	Gln	Arg	Val	Leu	Thr 270	Phe	Thr
Leu	Gly	Asp 275	Met	Gln	Pro	Gly	Glu 280	His	Arg	Thr	Ile	Thr 285	Val	Glu	Phe
Cys	Pro 290	Leu	Lys	Arg	Gly	Arg 295	Ala	Thr	Asn	Ile	Ala 300	Thr	Val	Ser	Tyr
Cys 305	Gly	Gly	His	Lys	Asn 310	Thr	Ala	Ser	Val	Thr 315	Thr	Val	Ile	Asn	Glu 320
Pro	Cys	Val	Gln	Val 325	Ser	Ile	Ala	Gly	Ala 330	Asp	Trp	Ser	Tyr	Val 335	Cys
Lys	Pro	Val	Glu 340	Tyr	Val	Ile	Ser	Val 345	Ser	Asn	Pro	Gly	Asp 350	Leu	Val
Leu	Arg	Asp 355	Val	Val	Val	Glu	Asp 360	Thr	Leu	Ser	Pro	Gly 365	Val	Thr	Val
Leu	Glu 370	Ala	Ala	Gly	Ala	Gln 375	Ile	Ser	Суз	Asn	180 380	Val	Val	Trp	Thr
Val 385	Lys	Glu	Leu	Asn	Pro 390	Gly	Glu	Ser	Leu	Gln 395	Tyr	Lys	Val	Leu	Val 400
Arg	Ala	Gln	Thr	Pro 405	Gly	Gln	Phe	Thr	Asn 410	Asn	Val	Val	Val	Lys 415	Ser
Суѕ	Ser	Asp	Cys 420	Gly	Thr	Cys	Thr	Ser 425	Суѕ	Ala	Glu	Ala	Thr 430	Thr	Tyr
ттр	Lys	Gly	Val	Ala	Ala	Thr	His	Met	Суз	Val	Val	Asp	Thr	Суѕ	Asp

440 435 445 Pro Val Cys Val Gly Glu Asn Thr Val Tyr Arg Ile Cys Val Thr Asn Arg Gly Ser Ala Glu Asp Thr Asn Val Ser Leu Met Leu Lys Phe Ser Lys Glu Leu Gln Pro Val Ser Phe Ser Gly Pro Thr Lys Gly Thr Ile 485 Thr Gly Asn Thr Val Val Phe Asp Ser Leu Pro Arg Leu Gly Ser Lys Glu Thr Val Glu Phe Ser Val Thr Leu Lys Ala Val Ser Ala Gly Asp Ala Arg Gly Glu Ala Ile Leu Ser Ser Asp Thr Leu Thr Val Pro Val Ser Asp Thr Glu Asn Thr His Ile Tyr <210> 442 <211> 192 <212> PRT <213> Chlamydia trachomatis serovar D Met Pro Glu Gly Glu Met Met His Lys Leu Gln Asp Val Ile Asp Arg Lys Leu-Leu Asp Ser Arg Arg Ile Phe Phe Ser Glu Pro Val Thr Glu Lys Ser Ala Thr Glu Ala Ile Lys Lys Leu Trp Tyr Leu Glu Leu Thr Asn Pro Gly Gln Pro Ile Val Phe Val Ile Asn Ser Pro Gly Gly Ser Val Asp Ala Gly Phe Ala Val Trp Asp Gln Ile Lys Met Ile Ser Ser Pro Leu Thr Thr Val Val Thr Gly Leu Ala Ala Ser Met Gly Ser Val Leu Ser Leu Cys Ala Val Pro Gly Arg Arg Phe Ala Thr Pro His Ala Arg Ile Met Ile His Gln Pro Ser Ile Gly Gly Thr Ile Thr Gly Gln 120 Ala Thr Asp Leu Asp Ile His Ala Arg Glu Ile Leu Lys Thr Lys Ala Arg Ile Ile Asp Val Tyr Val Glu Ala Thr Gly Gln Ser Arg Glu Val Ile Glu Lys Ala Ile Asp Arg Asp Met Trp Met Ser Ala Asn Glu Ala

175 165 170 Met Glu Phe Gly Leu Leu Asp Gly Ile Leu Phe Ser Phe Asn Asp Leu 185 180 <210> 443 <211> 275 <212> PRT <213> Chlamydia trachomatis serovar D <400> 443 Met Gly Phe Ser Ser Leu Leu Thr Thr Cys Arg Tyr Leu Leu Tyr Ser Gly Ala Gly Asn Ser Phe Ile Leu Gly Glu Ser Met Pro Ser Leu Glu Asp Val Leu Phe Leu Cys Gln Glu Glu Met Val Asp Gly Phe Leu Cys Val Glu Ser Ser Glu Ile Ala Asp Ala Lys Leu Thr Val Phe Asn Ser Asp Gly Ser Ile Ala Ser Met Cys Gly Asn Gly Leu Arg Cys Ala Met Ala His Val Ala Gln Cys Phe Gly Leu Glu Asp Val Ser Ile Glu Thr Glu Arg Gly Val Tyr Gln Gly Lys Phe Phe Ser Met Asn Arg Val Leu Val Asp Met Thr Leu Pro Asp Trp Lys Lys Ala Glu Arg Lys Leu Thr His Val Leu Pro Gly Met Pro Glu Gln Val Phe Phe Ile Asp Thr Gly Val Pro His Val Val Val Phe Val Ser Asp Leu Ser Lys Val Pro Val 155 Gln Glu Trp Gly Ser Phe Leu Arg Tyr His Glu Asp Phe Ala Pro Glu Gly Val Asn Val Asp Phe Val Gln Arg Lys Lys Asp Asp Leu Leu Leu Val Tyr Thr Tyr Glu Arg Gly Cys Glu Arg Glu Thr Leu Ser Cys Gly Thr Gly Met Leu Ala Ser Ala Leu Val Ala Ala Asp Ile Phe Ser Leu Gly Gln Asp Phe Ser Ile Ala Val Cys Ser Arg Ser Arg Asn Leu Ile Lys Ile Phe Ser Glu Lys Gly Lys Val Phe Leu Glu Gly Pro Val Ser Leu Leu Asn Arg Ser Glu Asn Phe Gly Trp Leu Glu Pro Lys Ser Arg 244

260 265 270

Arg Phe Gly 275

<210> 444

<211> 1770

<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 444

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Val Thr Glu Ala Ser Ser Ile Gln Asp Gln Ile Lys Asn Thr Asp Cys 20 25 30

Asn Val Ser Lys Leu Gly Tyr Ser Thr Ser Gln Ala Phe Thr Asp Met 35 40

Met Leu Ala Asp Asn Thr Glu Tyr Arg Ala Ala Asp Ser Val Ser Phe 50 60

Tyr Asp Phe Ser Thr Ser Ser Arg Leu Pro Arg Lys His Leu Ser Ser 65 70 .75 80

Ser Ser Glu Ala Ser Pro Thr Thr Glu Gly Val Ser Ser Ser Ser Ser Ser 90 95

Ile Tyr Ala Arg Glu Lys Leu Thr Ile Ser Glu Ser Gln Asp Ser Leu 115 120 125

Ser Asn Gln Ser Ile Glu Leu His Asp Asn Ser Ile Phe Gly Glu 130 135 140

Gly Glu Val Ile Phe Asp His Arg Val Ala Leu Lys Asn Gly Gly Ala 145 150150155160

Ile Tyr Gly Glu Lys Glu Val Val Phe Glu Asn Ile Lys Ser Leu Leu 165 170 175

Val Glu Val Asn Ile Ala Val Glu Lys Gly Gly Ser Val Tyr Ala Lys 180 185 190

Glu Arg Val Ser Leu Glu Asn Val Thr Glu Ala Thr Phe Ser Ser Asn 195 200 205

Gly Gly Glu Gln Gly Gly Gly Ile Tyr Ser Glu Gln Asp Met Leu 210 215 220

Ile Ser Asp Cys Asn Asn Val His Phe Gln Gly Asn Ala Ala Gly Ala 225 230 235 240

Thr Ala Val Lys Gln Cys Leu Asp Glu Glu Met Ile Val Leu Leu Ala 245 250 255

Glu Cys Val Asp Ser Leu Ser Glu Asp Thr Leu Asp Ser Thr Pro Glu

			260					265					270		
Thr	Glu	Gln 275	Thr	Glu	Ser	Asn	Gly 280	Asn	Gln	Asp	Gly	Ser 285	Ser	Glu	Thr
Glu	Asp 290	Thr	Gln	Val	Ser	Glu 295	Ser	Pro	Glu	Ser	Thr 300	Pro	Ser	Pro	Asp
Asp 305	Val	Leu	Gly	Lys	Gly 310	Gly	Gly	Ile	Tyr	Thr 315	Glu	Lys	Ser	Leu	Thr 320
Ile	Thr	Gly	Ile	Thr 325	Gly	Thr	Ile	Asp	Phe 330	Val	Ser	Asn	Ile	Ala 335	Thr
Asp	Ser	Gly	Ala 340	Gly	Val	Phe	Thr	Lys 345	Glu	Asn	Leu	Ser	Cys 350	Thr	Asn
Thr	Asn	Ser 355	Leu	Gln	Phe	Leu	Lys 360	Asn	Ser	Ala	Gly	Gln 365	His	Gly	Gly
Gly	Ala 370	Tyr	Val	Thr	Gln	Thr 375	Met	Ser	Val	Thr	Asn 380	Thr	Thr	Ser	Glu
Ser 385	Ile	Thr	Thr	Pro	Pro 390	Leu	Ile	Gly	Glu	Val 395	Ile	Phe	Ser	Glu	Asn 400
Thr	Ala	Lys	Gly	His 405	Gly	Gly	Gly	Ile	Cys 410	Thr	Asn	Lys	Leu	Ser 415	Leu
Ser	Asn	Leu	Lys 420	Thr	Val	Thr	Leu	Thr 425	Lys	Asn	Ser	Ala	Lys 430	Glu	Ser
Gly	Gly	Ala 435	Ile	Phe	Thr	Asp	Leu 440	Ala	Ser	Ile	Pro	Ile 445	Thr	Asp	Thr
Pro	Glu 450	Ser	Ser	Thr	Pro	Ser 455	Ser	Ser	Ser	Pro	Ala 460	Ser	Thr	Pro	Glu
Val 465	Val	Ala	Ser	Ala	Lys 470	Ile	Asn	Arg	Phe	Phe 475	Ala	Ser	Thr	Ala	Lys 480
Pro	Ala	Ala	Pro	Ser 485	Leu	Thr	Glu	Ala	Glu 490	Ser	Asp	Gln	Thr	Asp 495	Gln
Thr	Glu	Thr	Ser 500	Asp	Thr	Asn	Ser	Asp 505	Ile	Asp	Val	Ser	Ile 510	Glu	Asn
Ile	Leu	Asn 515	Val	Ala	Ile	Asn	Gln 520	Asn	Thr	Ser	Ala	Lys 525	Lys	Gly	Gly
Ala	Ile 530	Tyr	Gly	Lys	Lys	Ala 535	Lys	Leu	Ser	Arg	Ile 540	Asn	Asn	Leu	Glu
Leu 545	Ser	Gly	Asn	Ser	Ser 550	Gln	Asp	Val	Gly	Gly 555	Gly	Leu	Cys	Leu	Thr 560
Glu	Ser	Val	Glu	Phe 565	Asp	Ala	Ile	Gly	Ser 570	Leu	Leu	Ser	His	Tyr 575	Asn
Ser	Ala	Ala	Lys 580	Glu	Gly	Gly	Ala	Ile 585	His	Ser	Lys	Thr	Val 590	Thr	Leu

Ser	Asn	Leu 595	Lys	Ser	Thr	Phe	Thr 600	Phe	Ala	Asp	Asn	Thr 605	Val	Lys	Ala
Ile	Val 610	Glu	Ser	Thr	Pro	Glu 615	Ala	Pro	Glu	Glu	Ile 620	Pro	Pro	Val	Glu
Gly 625	Glu	Glu	Ser	Thr	Ala 630	Thr	Glu	Asp	Pro	Asn 635	Ser	Asn	Thr	Glu	Gly 640
Ser	Ser	Ala	Asn	Thr 645	Asn	Leu	Glu	Gly	Ser 650	Gln	Gly	Asp	Thr	Ala 655	Asp
Thr	Gly	Thr	Gly 660	Asp	Val	Asn	Asn	Glu 665	Ser	Gln	Asp	Thr	Ser 670	Asp	Thr
Gly	Asn	Ala 675	Glu	Ser	Glu	Glu	Gln 680	Leu	Gln	Asp	Ser	Thr 685	Gln	Ser	Asn
Glu	Glu 690	Asn	Thr	Leu	Pro	Asn 695	Ser	Asn	Ile	Asp	Gln 700	Ser	Asn	Glu	Asn
Thr 705	Asp	Glu	Ser	Ser	Asp 710	Ser	His	Thr	Glu	Glu 715	Ile	Thr	Asp	Glu	Ser 720
Val	Ser	Ser	Ser	Ser 725	Glu	Ser	Gly	Ser	Ser 730	Thr	Pro	Gln	Asp	Gly 735	Gly
Ala	Ala	Ser	Ser 740	Gly	Ala	Pro	Ser	Gly 745	Asp	G1n	Ser	Ile	Ser 750	Ala	Asn
Ala	Cys	Leu 755	Ala	Lys	Ser	Tyr	Ala 760	Ala	Ser	Thr	Asp	Ser 765	Ser	Pro	Val
Ser	Asn 770	Ser	Ser	Gly	Ser	Glu 775	Glu	Pro	Val	Thr	Ser 780	Ser	Ser	Asp	Ser
Asp 785	Val	Thr	Ala	Ser	Ser 790	Asp	Asn	Pro	Asp	Ser 795	Ser	Ser	Ser	Gly	Asp 800
Ser	Ala	Gly	Asp	Ser 805	Glu	Glu	Pro	Thr	Glu 810	Pro	Glu	Ala	Gly	Ser 815	Thr
Thr	Glu	Thr	Leu 820	Thr	Leu	Ile	Gly	Gly 825	Gly	Ala	Ile	Tyr	Gly 830	Glu	Thr
Val	Lys	11e 835	Glu	Asn	Phe	Ser	Gly 840	Gln	Gly	Ile	Phe	Ser 845	Gly	Asn	Lys
Ala	Ile 850	Asp	Asn	Thr	Thr	Glu 855	Gly	Ser	Ser	Ser	Lys 860	Ser	Asp	Val	Leu
Gly 865	Gly	Ala	Val '	Tyr	Ala 870	Lys	Thr	Leu	Phe	Asn 875	Leu	Asp	Ser	Gly	Ser 880
Ser	Arg	Arg	Thr	Val 885	Thr	Phe	Ser	Gly	Asn 890	Thr	Val	Ser	Ser	Gln 895	Ser
Thr	Thr	Gly	Gln 900	Val	Ala	Gly	Gly	Ala 905	Ile	Tyr	Ser	Pro	Thr 910	Val	Thr

- Ile Ala Thr Pro Val Val Phe Ser Lys Asn Ser Ala Thr Asn Asn Ala 915 920 925
- Asn Asn Thr Thr Asp Thr Gln Arg Lys Asp Thr Phe Gly Gly Ala Ile 930 940
- Gly Ala Thr Ser Ala Val Ser Leu Ser Gly Gly Ala His Phe Leu Glu 945 950 955 960
- Asn Val Ala Asp Leu Gly Ser Ala Ile Gly Leu Val Pro Gly Thr Gln $965 \\ 970 \\ 975$
- Asn Thr Glu Thr Val Lys Leu Glu Ser Gly Ser Tyr Tyr Phe Glu Lys 980 985 990
- Asn Lys Ala Leu Lys Arg Ala Thr Ile Tyr Ala Pro Val Val Ser Ile 995 1000 1005
- Lys Ala Tyr Thr Ala Thr Phe Asn Gln Asn Arg Ser Leu Glu Glu Gly 1010 1015 1020
- Ser Ala Ile Tyr Phe Thr Lys Glu Ala Ser Ile Glu Ser Leu Gly Ser 1025 1030 1035 1040
- Val Leu Phe Thr Gly Asn Leu Val Thr Leu Thr Leu Ser Thr Thr Thr 1045 1050 1055
- Glu Gly Thr Pro Ala Thr Thr Ser Gly Asp Val Thr Lys Tyr Gly Ala 1060 1065 1070
- Ala Ile Phe Gly Gln Ile Ala Ser Ser Asn Gly Ser Gln Thr Asp Asn 1075 1080 1085
- Leu Pro Leu Lys Leu Ile Ala Ser Gly Gly Asn Ile Cys Phe Arg Asn 1090 1095 1100
- Asn Glu Tyr Arg Pro Thr Ser Ser Asp Thr Gly Thr Ser Thr Phe Cys 1105 1110 1115 1120
- Ser Ile Ala Gly Asp Val Lys Leu Thr Met Gln Ala Ala Lys Gly Lys 1125 1130 1135
- Thr Ile Ser Phe Phe Asp Ala Ile Arg Thr Ser Thr Lys Lys Thr Gly 1140 1145 1150
- Thr Gln Ala Thr Ala Tyr Asp Thr Leu Asp Ile Asn Lys Ser Glu Asp 1155 1160 1165
- Ser Glu Thr Val Asn Ser Ala Phe Thr Gly Thr Ile Leu Phe Ser Ser 1170 1175 1180
- Glu Leu His Glu Asn Lys Ser Tyr Ile Pro Gln Asn Val Val Leu His 1185 1190 1195 1200
- Ser Gly Ser Leu Val Leu Lys Pro Asn Thr Glu Leu His Val Ile Ser 1205 1210 1215
- Phe Glu Gln Lys Glu Gly Ser Ser Leu Val Met Thr Pro Gly Ser Val 1220 1225 1230
- Leu Ser Asn Gln Thr Val Ala Asp Gly Ala Leu Val Ile Asn Asn Met

		1235					1240)				1245	,		
Thr	Ile 1250	_	Leu	Ser	Ser	Val 1255		Lys	Asn	Gly	Ile 1260		Glu	Gly	Asn
Ile 1265	Phe	Thr	Pro	Pro	Glu 1270		Arg	Ile	Ile	Asp 1275		Thr	Thr	Gly	Gly 1280
Ser	Gly	Gly.	Thr	Pro 1285		Thr	Asp	Ser	Glu 1290		Asn	Gln	Asn	Ser 1295	
Asp	Thr	Glu	Glu 1300		Asn	Asn	Asn	Asp 1305		Ser	Asn	Gln	Gly 1310		Ser
Ala	Asn	Gly 1315		Ser	Ser	Pro	Ala 1320		Ala	Ala	Ala	His 1325		Ser	Arg
Thr	Arg 1330		Phe	Ala	Ala	Ala 1335		Thr	Ala	Thr	Pro 1340		Thr	Thr	Pro
Thr 1345	Ala	Thr	Thr	Thr	Thr 1350		Asn	Gln	Val	Ile 1355		Gly	Gly	Glu	Ile 1360
Lys	Leu	Ile	Asp	Pro 1365		Gly	Thr	Phe	Phe 1370		Asn	Pro	Ala	Leu 1375	
Ser	Asp	Gln	Gln 1380		Ser	Leu	Leu	Val 1385		Pro	Thr	Asp	Ser 1390		Lys
Met	Gln	Ala 1395		Lys	Ile	Val	Leu 1400		Gly	Asp	Ile	Ala 1405		Gln	Lys
Gly	Tyr 1410		Gly	Thr	Leu	Thr 1415		Asp	Pro	Asp	Gln 1420		Gln	Asn	Gly
Thr 1425	Ile	Ser	Val	Leu	Trp 1430		Phe	Asp	Ser	Tyr 1435		Gln	Trp	Ala	Tyr 1440
Val	Pro	Arg	Asp	Asn 1445		Phe	Tyr	Ala	Asn 1450		Ile	Leu	Gly	Ser 1455	
Met	Leu	Met	Val 1460		Val	Lys	Gln	Gly 1465		Leu	Asn	Asp	Lys 1470		Asn
Leu	Ala	Arg 1475		Glu	Glu	Val	Ser 1480		Asn	Asn	Leu	Trp 1485		Ser	Gly
Leu	Gly 1490		Met	Leu	Ser	Gln 1495		Gly	Thr	Pro	Thr 1500		Glu	Glu	Phe
Thr 1505	Tyr	Tyr	Ser	Arg	Gly 1510		Ser	Val	Ala	Leu 1515		Ala	Lys	Pro	Ala 1520
His	Asp	Val	Ile	Val 1525		Ala	Ala	Phe	Ser 1530		Met	Ile	Gly	Lys 1535	
Lys	Ser	Leu	Lys 1540		Glu	Asn	Asn	Tyr 1545		His	Lys	Gly	Ser 1550		Tyr
Ser	Tyr	Gln 1555		Ser	Val	Tyr	Gly 1560		Lys	Pro	Phe	His 1565		Val	Ile

Asn Lys Lys Thr Glu Lys Ser Leu Pro Leu Leu Leu Gln Gly Val Ile Ser Tyr Gly Tyr Ile Lys His Asp Thr Val Thr His Tyr Pro Thr Ile 1590 Arg Glu Arg Asn Lys Gly Glu Trp Glu Asp Leu Gly Trp Leu Thr Ala 1610 Leu Arg Val Ser Ser Val Leu Arg Thr Pro Ala Gln Gly Asp Thr Lys 1625 Arg Ile Thr Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln Phe Thr Glu Thr Glu Tyr Asp Pro Arg Tyr Phe Asp Asn Cys Thr 1655 1660 Tyr Arg Asn Leu Ala Ile Pro Met Gly Leu Ala Phe Glu Gly Glu Leu Ser Gly Asn Asp Ile Leu Met Tyr Asn Arg Phe Ser Val Ala Tyr Met 1690 Leu Ser Ile Tyr Arg Asn Ser Pro Thr Cys Lys Tyr Gln Val Leu Ser 1705 Ser Gly Glu Gly Glu Ile Ile Cys Gly Val Pro Thr Arg Asn Ser Ala Arg Gly Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Leu Trp Thr Leu Tyr Gly Ser Tyr Thr Ile Glu Ala Asp Ala His Thr Leu Ala His 1755 Met Met Asn Cys Gly Ala Arg Met Thr Phe 1765

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Val Ser Gly Phe Cys Phe Pro Glu Pro Lys Glu Leu Asn Phe Ser Arg
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Val Gly Thr Ser Ser Ser Thr Thr Phe Thr Glu Thr Val Gly Glu Ala 35 40 45

Gly Ala Glu Tyr Ile Val Ser Gly Asn Ala Ser Phe Thr Lys Phe Thr 50 55 60

Asn Ile Pro Thr Thr Asp Thr Thr Thr Pro Thr Asn Ser Asn Ser Ser 65 70 75 80

Ser	Ser	Asn	Gly	Glu 85	Thr	Ala	Ser	Val	Ser 90	Glu	Asp	Ser	Asp	Ser 95	Thr
Thr	Thr	Thr	Pro 100	Asp	Pro	Lys	Gly	Gly 105	Gly	Ala	Phe	Tyr	Asn 110	Ala	His
Ser	Gly	Val 115	Leu	Ser	Phe	Met	Thr 120	Arg	Ser	Gly	Thr	Glu 125	Gly	Ser	Leu
Thr	Leu 130	Ser	Glu	Ile	Lys	Ile 135	Thr	Gly	Glu	Gly	Gly 140	Ala	Ile	Phe	Ser
Gln 145	Gly	Glu	Leu	Leu	Phe 150	Thr	Asp	Leu	Thr	Gly 155	Leu	Thr	Ile	Gln	Asn 160
Asn	Leu	Ser	Gln	Leu 165	Ser	Gly	Gly	Ala	Ile 170	Phe	Gly	Glu	Ser	Thr 175	Ile
Ser	Leu	Ser	Gly 180	Ile	Thr	Lys	Ala	Thr 185	Phe	Ser	Ser	Asn	Ser 190	Ala	Glu
Val	Pro	Ala 195	Pro	Val	Lys	Lys	Pro 200	Thr	Glu	Pro	Lys	Ala 205	Gln	Thr	Ala
Ser	Glu 210	Thr	Ser	Gly	Ser	Ser 215	Ser	Ser	Ser	Gly	Asn 220	Asp	Ser	Val	Ser
Ser 225	Pro	Ser	Ser	Ser	Arg 230	Ala	Glu	Pro	Ala	Ala 235	Ala	Asn	Leu	Gln	Ser 240
His	Phe	Ile	Суѕ	Ala 245	Thr	Ala	Thr	Pro	Ala 250	Ala	Gln	Thr	Asp	Thr 255	Glu
Thr	Ser	Thr	Pro 260	Ser	His	Lys	Pro	Gly 265	Ser	Gly	Gly	Ala	Ile 270	Tyr	Ala
Lys	Gly	Asp 275	Leu	Thr	Ile	Ala	Asp 280	Ser	Gln	Glu	Val	Leu 285	Phe	Ser	Ile
Asn	Lys 290	Ala	Thr	Lys	Asp	Gly 295	Gly	Ala	Ile	Phe	Ala 300	Glu	Lys	Asp	Val
Ser 305	Phe	Glu	Asn	Ile	Thr 310	Ser	Leu	Lys	Val	Gln 315	Thr	Asn	Gly	Ala	Glu 320
Glu	Lys	Gly	Gly	Ala 325	Ile	Tyr	Ala	Lys	Gly 330	Asp	Leu	Ser	Ile	Gln 335	Ser
Ser	Lys	Gln	Ser 340	Leu	Phe	Asn	Ser	Asn 345	Tyr	Ser	Lys	Gln	Gly 350	Gly	Gly
Ala	Leu	Tyr 355	Val	Glu	Gly	Asp	Ile 360	Asn	Phe	Gln	Asp	Leu 365	Glu	Glu	Ile
Arg	Ile 370	Lys	Tyr	Asn	Lys	Ala 375	Gly	Thr	Phe	Glu	Thr 380	Lys	Lys	Ile	Thr
Leu 385		Lys	Ala	Gln	Ala 390	Ser	Ala	Gly	Asn	Ala 395	Asp	Ala	Trp	Ala	Ser 400

PCT/US01/23121

Ser Ser Pro Gln Ser Gly Ser Gly Ala Thr Thr Val Ser Asn Ser Gly Asp Ser Ser Ser Gly Ser Asp Ser Asp Thr Ser Glu Thr Val Pro Ala Thr Ala Lys Gly Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser Ile Thr Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gln Gly Gly Gly Ile 490 Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asn Leu Thr Gly Lys Thr Leu Phe Gln Glu Asn Thr Ala Lys Glu Glu Gly Gly Leu Phe Ile Lys Gly Thr Asp Lys Ala Leu Thr Met Thr Gly Leu Asp Ser Phe Cys Leu Ile Asn Asn Thr Ser Glu Lys His Gly Gly Gly Ala Phe Val Thr Lys Glu Ile Ser Gln Thr Tyr Thr Ser Asp Val Glu Thr Ile Pro Gly Ile Thr Pro Val His Gly Glu Thr Val Ile Thr Gly Asn Lys Ser Thr Gly Gly Asn Gly Gly Val Cys Thr Lys Arg Leu Ala Leu Ser Asn Leu Gln Ser Ile Ser Ile Ser Gly Asn Ser Ala Ala Glu Asn Gly Gly Ala His Thr Cys Pro Asp Ser Phe Pro Thr Ala Asp Thr Ala Glu Gln Pro Ala Ala Ala Ser Ala Ala Thr Ser Thr Pro Glu Ser Ala Pro Val Val Ser Thr Ala Leu Ser Thr Pro Ser Ser Ser Thr Val Ser Ser Leu 665 Thr Leu Leu Ala Ala Ser Ser Gln Ala Ser Pro Ala Thr Ser Asn Lys 680 Glu Thr Gln Asp Pro Asn Ala Asp Thr Asp Leu Leu Ile Asp Tyr Val Val Asp Thr Thr Ile Ser Lys Asn Thr Ala Lys Lys Gly Gly Ile Tyr Ala Lys Lys Ala Lys Met Ser Arg Ile Asp Gln Leu Asn Ile Ser

	725	730	735
Glu Asn Ser Ala 740		ly Gly Ile Cys Cys	Lys Glu Ser 750
Leu Glu Leu Asp	Ala Leu Val Ser L	eu Ser Val Thr Glu	Asn Leu Val
755	760	765	
Gly Lys Glu Gly 770	Gly Gly Leu His A	ala Lys Thr Val Asn 780	Ile Ser Asn
Leu Lys Ser Gly	Phe Ser Phe Ser A	asn Asn Lys Ala Asn	Ser Ser Ser
785		795	800
Thr Gly Val Ala	Thr Thr Ala Ser A	ala Pro Ala Ala Ala	Ala Ala Ser
	805	810	815
Leu Gln Ala Ala 820		Pro Ser Ser Pro Ala 25	Thr Pro Thr 830 ·
Tyr Ser Gly Val	Val Gly Gly Ala I	le Tyr Gly Glu Lys	Val Thr Phe
835	840	845	
Ser Gln Cys Ser	Gly Thr Cys Gln F	Phe Ser Gly Asn Gln	Ala Ile Asp
850	855	860	
Asn Asn Pro Ser	Gln Ser Ser Leu A	sn Val Gln Gly Gly	Ala Ile Tyr
865	870	875	880
Ala Lys Thr Ser	Leu Ser Ile Gly S	Ser Ser Asp Ala Gly	Thr Ser Tyr
	885	890	895
Ile Phe Ser Gly		hr Gly Lys Ser Gln	Thr Thr Gly
900		105	910
Gln Ile Ala Gly	Gly Ala Ile Tyr S	Ser Pro Thr Val Thr	Leu Asn Cys
915	920	925	
Pro Ala Thr Phe 930	Ser Asn Asn Thr A	ala Ser Met Ala Thr 940	Pro Lys Thr
Ser Ser Glu Asp	Gly Ser Ser Gly A	sn Ser Ile Lys Asp	Thr Ile Gly
945	950	955	960
Gly Ala Ile Ala	Gly Thr Ala Ile T	hr Leu Ser Gly Val	Ser Arg Phe
	965	970	975
Ser Gly Asn Thr		ala Ala Ile Gly Thr	Leu Ala Asn
980		185	990
Ala Asn Thr Pro 995	Ser Ala Thr Ser 6	Gly Ser Gln Asn Ser 1005	
Lys Ile Thr Leu	Glu Asn Gly Ser F	Phe Ile Phe Glu Arg	Asn Gln Ala
1010	1015	1020	
Asn Lys Arg Gly	Ala Ile Tyr Ser F	Pro Ser Val Ser Ile	Lys Gly Asn
1025	1030	1035	1040
Asn Ile Thr Phe	Asn Gln Asn Thr S	er Thr His Asp Gly	Ser Ala Ile
	1045	1050	1055

Tyr Phe Thr Lys Asp Ala Thr Ile Glu Ser Leu Gly Ser Val Leu Phe 1060 1065 1070

- Thr Gly Asn Asn Val Thr Ala Thr Gln Ala Ser Ser Ala Thr Ser Gly 1075 1080 1085
- Gln Asn Thr Asn Thr Ala Asn Tyr Gly Ala Ala Ile Phe Gly Asp Pro 1090 1095 1100
- Gly Thr Thr Gln Ser Ser Gln Thr Asp Ala Ile Leu Thr Leu Leu Ala 1105 1110 1115 1120
- Ser Ser Gly Asn Ile Thr Phe Ser Asn Asn Ser Leu Gln Asn Asn Gln 1125 1130 1135
- Gly Asp Thr Pro Ala Ser Lys Phe Cys Ser Ile Ala Gly Tyr Val Lys 1140 1145 1150
- Leu Ser Leu Gln Ala Ala Lys Gly Lys Thr Ile Ser Phe Phe Asp Cys 1155 1160 1165
- Val His Thr Ser Thr Lys Lys Ile Gly Ser Thr Gln Asn Val Tyr Glu 1170 1175 1180
- Thr Leu Asp Ile Asn Lys Glu Glu Asn Ser Asn Pro Tyr Thr Gly Thr 1185 1190 1195 1200
- Ile Val Phe Ser Ser Glu Leu His Glu Asn Lys Ser Tyr Ile Pro Gln 1205 1210 1215
- Asn Ala Ile Leu His Asn Gly Thr Leu Val Leu Lys Glu Lys Thr Glu 1220 1225 1230
- Leu His Val Val Ser Phe Glu Gln Lys Glu Gly Ser Lys Leu Ile Met 1235 1240 1245
- Lys Pro Gly Ala Val Leu Ser Asn Gln Asn Ile Ala Asn Gly Ala Leu 1250 1255 1260
- Val Ile Asn Gly Leu Thr Ile Asp Leu Ser Ser Met Gly Thr Pro Gln 1265 1270 1275 1280
- Ala Gly Glu Ile Phe Ser Pro Pro Glu Leu Arg Ile Val Ala Thr Thr 1285 1290 1295
- Ser Ser Ala Ser Gly Gly Ser Gly Val Ser Ser Ser Ile Pro Thr Asn 1300 1305 1310
- Pro Lys Arg Ile Ser Ala Ala Ala Pro Ser Gly Ser Ala Ala Thr Thr 1315 1320 1325
- Pro Thr Met Ser Glu Asn Lys Val Phe Leu Thr Gly Asp Leu Thr Leu 1330 1335 1340
- Ile Asp Pro Asn Gly Asn Phe Tyr Gln Asn Pro Met Leu Gly Ser Asp 1345 1350 1355 1360
- Leu Asp Val Pro Leu Ile Lys Leu Pro Thr Asn Thr Ser Asp Val Gln
 1365 1370 1375

Val Tyr Asp Leu Thr Leu Ser Gly Asp Leu Phe Pro Gln Lys Gly Tyr 1380 1385 1390

- Met Gly Thr Trp Thr Leu Asp Ser Asn Pro Gln Thr Gly Lys Leu Gln 1395 1400 1405
- Ala Arg Trp Thr Phe Asp Thr Tyr Arg Arg Trp Val Tyr Ile Pro Arg 1410 1415 1420
- Asp Asn His Phe Tyr Ala Asn Ser Ile Leu Gly Ser Gln Asn Ser Met 1425 1430 1435 1440
- Ile Val Val Lys Gln Gly Leu Ile Asn Asn Met Leu Asn Asn Ala Arg
 1445 1450 1455
- Phe Asp Asp Ile Ala Tyr Asn Asn Phe Trp Val Ser Gly Val Gly Thr 1460 1465 1470
- Phe Leu Ala Gln Gln Gly Thr Pro Leu Ser Glu Glu Phe Ser Tyr Tyr 1475 1480 1485
- Ser Arg Gly Thr Ser Val Ala Ile Asp Ala Lys Pro Arg Gln Asp Phe 1490 1495 1500
- Ile Leu Gly Ala Ala Phe Ser Lys Met Val Gly Lys Thr Lys Ala Ile 1505 1510 1515 1520
- Lys Lys Met His Asn Tyr Phe His Lys Gly Ser Glu Tyr Ser Tyr Gln 1525 1530 1535
- Ala Ser Val Tyr Gly Gly Lys Phe Leu Tyr Phe Leu Leu Asn Lys Gln 1540 . 1545 . 1550
- His Gly Trp Ala Leu Pro Phe Leu Ile Gln Gly Val Val Ser Tyr Gly 1555 1560 1565
- His Ile Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu Arg 1570 1575 1580
- Asn Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg Ile 1585 1590 1595 1600
- Ser Met Asp Leu Lys Glu Pro Ser Lys Asp Ser Ser Lys Arg Ile Thr 1605 1610 1615
- Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln Phe Thr 1620 1625 1630
- Glu Ile Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala Tyr Arg Asn 1635 1640 1645
- Leu Ser Leu Pro Val Gly Cys Ala Val Glu Gly Ala Ile Met Asn Cys 1650 1655 1660
- Asn Ile Leu Met Tyr Asn Lys Leu Ala Leu Ala Tyr Met Pro Ser Ile 1665 1670 1680
- Tyr Arg Asn Asn Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asn Glu 1685 1690 1695
- Ala Gly Gln Val Ile Cys Gly Val Pro Thr Arg Thr Ser Ala Arg'Ala

255

1700 1705 1710

Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr Gly 1715 1720 1725

Asn Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gln Met Thr Ser 1730 1735 1740

Cys Gly Ala Arg Met Ile Phe 1745 1750

<210> 446

<211> 660

<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 446

Met Ser Glu Lys Arg Lys Ser Asn Lys Ile Ile Gly Ile Asp Leu Gly

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15

Thr Thr Asn Ser Cys Val Ser Val Met Glu Gly Gly Gln Pro Lys Val 20 25 30

Ile Ala Ser Ser Glu Gly Thr Arg Thr Thr Pro Ser Ile Val Ala Phe 35 40

Lys Gly Glu Thr Leu Val Gly Ile Pro Ala Lys Arg Gln Ala Val 50 55 60

Thr Asn Pro Glu Lys Thr Leu Ala Ser Thr Lys Arg Phe Ile Gly Arg 65 70 75 80

Lys Phe Ser Glu Val Glu Ser Glu Ile Lys Thr Val Pro Tyr Lys Val 85 90 95

Ala Pro Asn Ser Lys Gly Asp Ala Val Phe Asp Val Glu Gln Lys Leu 100 105 110

Tyr Thr Pro Glu Glu Ile Gly Ala Gln Ile Leu Met Lys Met Lys Glu 115 120 125

Thr Ala Glu Ala Tyr Leu Gly Glu Thr Val Thr Glu Ala Val Ile Thr 130 140

Val Pro Ala Tyr Phe Asn Asp Ser Gln Arg Ala Ser Thr Lys Asp Ala 145 150 155 160

Gly Arg Ile Ala Gly Leu Asp Val Lys Arg Ile Ile Pro Glu Pro Thr 165 170 175

Ala Ala Leu Ala Tyr Gly Ile Asp Lys Glu Gly Asp Lys Ile 180 185 190

Ala Val Phe Asp Leu Gly Gly Thr Phe Asp Ile Ser Ile Leu Glu 195 200 205

Ile Gly Asp Gly Val Phe Glu Val Leu Ser Thr Asn Gly Asp Thr His 210 225 220

Leu Gly Gly Asp Asp Phe Asp Gly Val Ile Ile Asn Trp Met Leu Asp

WO 02/08267

256

225					230					235					240
Glu	Phe	Lys	Lys	Gln 245	Glu	Gly	Ile	Asp	Leu 250	Ser	Lys	Asp	Asn	Met 255	Ala
Leu	Gln	Arg	Leu 260	Lys	Asp	Ala	Ala	Glu 265	Lys	Ala	Lys	Ile	Glu 270	Leu	Ser
Gly	Val	Ser 275	Ser	Thr	Glu	Ile	Asn 280	Gln	Pro	Phe	Ile	Thr 285	Ile	Asp	Ala
Asn	Gly 290	Pro	Lys	His	Leu	Ala 295	Leu	Thr	Leu	Thr	Arg 300	Ala	Gln	Phe	Glu
His 305	Leu	Ala	Ser	Ser	Leu 310	Ile	Glu	Arg	Thr	Lys 315	Gln	Pro	Суз	Ala	Gln 320
Ala	Leu	Lys	Asp	Ala 325	Lys	Leu	Ser	Ala	Ser 330	Asp	Ile	Asp	Asp	Val 335	Leu
Leu	Val	Gly	Gly 340	Met	Ser	Arg	Met	Pro 345	Ala	Val	Gln	Ala	Val 350	Val	Lys
Glu	Ile	Phe 355	Gly	Lys	Glu	Pro	Asn 360	Lys	Gly	Val	Asn	Pro 365	Asp	Glu	Val
Val	Ala 370	Ile	Gly	Ala	Ala	Ile 375	Gln	Gly	Gly	Val	Leu 380	Gly	Gly	Glu	Val
Lys 385	Asp	Val	Leu	Leu	Leu 390	Asp	Val	Ile	Pro	Leu 395	Ser	Leu	Gly	Ile	Glu 400
Thr	Leu	Gly	Gly	Val 405	Met	Thr	Pro	Leu	Val 410	Glu	Arg	Asn	Thr	Thr 415	Ile
Pro	Thr	Gln	Lys 420	Lys	Gln	Ile	Phe	Ser 425	Thr	Ala	Ala	Asp	Asn 430	Gln	Pro
Ala	Val	Thr 435	Ile	Val	Val	Leu	Gln 440	Gly	Glu	Arg	Pro	Met 445	Ala	Lys	Asp
Asn	Lys 450	Glu	Ile	Gly	Arg	Phe 455	Asp	Leu	Thr	Asp	Ile 460	Pro	Pro	Ala	Pro
Arg 465	Gly	His	Pro	Gln	Ile 470	Glu	Val	Thr	Phe	Asp 475	Ile	Asp	Ala	Asn	Gly 480
Ile	Leu	His	Val	Ser 485	Ala	Lys	Asp	Ala	Ala 490	Ser	Gly	Arg	Glu	Gln 495	Lys
Ile	Arg	Ile	Glu 500	Ala	Ser	Ser	Gly	Leu 505	Lys	Glu	Asp	Glu	Ile 510	Gln	Gln
Met	Ile	Arg 515	Asp	Ala	Glu	Leu	His 520	Lys	Glu	Glu	Asp	Lys 525	Gln	Arg	Lys
Glu	Ala 530	Ser	Asp	Val	Lys	Asn 535	Glu	Ala	Asp	Gly	Met 540	Ile	Phe	Arg	Ala
Glu 545	Lys	Ala	Val	Lys	Asp 550	Tyr	His	Asp	Lys	Ile 555	Pro	Ala	Glu	Leu	Val 560

Lys Glu Ile Glu Glu His Ile Glu Lys Val Arg Gln Ala Ile Lys Glu

Asp Ala Ser Thr Thr Ala Ile Lys Ala Ala Ser Asp Glu Leu Ser Thr

His Met Gln Lys Ile Gly Glu Ala Met Gln Ala Gln Ser Ala Ser Ala

Ala Ala Ser Ser Ala Ala Asn Ala Gln Gly Gly Pro Asn Ile Asn Ser 615

Glu Asp Leu Lys Lys His Ser Phe Ser Thr Arg Pro Pro Ala Gly Gly

Ser Ala Ser Ser Thr Asp Asn Ile Glu Asp Ala Asp Val Glu Ile Val

Asp Lys Pro Glu 660

<210> 447

<211> 326

<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 447

Met Val Ser Gln Thr Val Ser Val Ala Val Thr Gly Gly Thr Gly Gln

Ile Ala Tyr Ser Phe Leu Phe Ser Leu Ala His Gly Asp Val Phe Gly

Leu Asp Cys Gly Ile Asp Leu Arg Ile Tyr Asp Ile Pro Gly Thr Glu

Arg Ala Leu Ser Gly Val Arg Met Glu Leu Asp Asp Gly Ala Phe Pro

Leu Leu Gln Arg Val Gln Val Thr Thr Ser Leu His Asp Ala Phe Asp

Gly Ile Asp Ala Ala Phe Leu Ile Gly Ser Val Pro Arg Gly Pro Gly

Met Glu Arg Arg Asp Leu Leu Lys Lys Asn Gly Glu Ile Phe Ala Thr

Gln Gly Lys Ala Leu Asn Thr Thr Ala Lys Arg Asp Ala Lys Ile Phe

Val Val Gly Asn Pro Val Asn Thr Asn Cys Trp Ile Ala Met Asn His

Ala Pro Arg Leu Arg Lys Asn Phe His Ala Met Leu Arg Leu Asp

Gln Asn Arg Met His Ser Met Leu Ser His Arg Ala Glu Val Pro Leu

258

Ser Ala Val Ser Gln Val Val Val Trp Gly Asn His Ser Ala Lys Gln 180 185 190

Val Pro Asp Phe Thr Gln Ala Leu Ile Asn Asp Arg Pro Ile Ala Glu 195 200 205

Thr Ile Ala Asp Arg Asp Trp Leu Glu Asn Ile Met Val Pro Ser Val 210 220

Gln Ser Arg Gly Ser Ala Val Ile Glu Ala Arg Gly Lys Ser Ser Ala 225 230 235 240

Ala Ser Ala Ala Arg Ala Leu Ala Glu Ala Ala Arg Ser Ile Tyr Gln 245 250 255

Pro Lys Glu Gly Glu Trp Phe Ser Ser Gly Val Cys Ser Asp His Asn 260 265 270

Pro Tyr Gly Leu Pro Glu Asp Leu Ile Phe Gly Phe Pro Cys Arg Met 275 280 285

Leu Ala Thr Gly Glu Tyr Glu Val Ile Pro Arg Leu Pro Trp Asp Ala 290 295 300

Phe Ile Arg Gly Lys Met Gln Ile Ser Leu Asp Glu Ile Leu Gln Glu 305 310 315 320

Lys Ala Ser Val Ser Leu 325

<210> 448

<211> 232

<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 448

Met Thr Lys His Gly Lys Arg Ile Arg Gly Ile Gln Glu Thr Tyr Asp
5 10 15

Leu Ala Lys Ser Tyr Ser Leu Gly Glu Ala Ile Asp Ile Leu Lys Gln
20 25 30

Cys Pro Thr Val Arg Phe Asp Gln Thr Val Asp Val Ser Val Lys Leu 35 40 45

Gly Ile Asp Pro Arg Lys Ser Asp Gln Gln Ile Arg Gly Ser Val Ser 50 60

Leu Pro His Gly Thr Gly Lys Val Leu Arg Ile Leu Val Phe Ala Ala 65 70 75 80

Gly Asp Lys Ala Ala Glu Ala Ile Glu Ala Gly Ala Asp Phe Val Gly 85 90 95

Ser Asp Asp Leu Val Glu Lys Ile Lys Gly Gly Trp Val Asp Phe Asp 100 105 110

Val Ala Val Ala Thr Pro Asp Met Met Arg Glu Val Gly Lys Leu Gly
115 120 125

Lys Val Leu Gly Pro Arg Asn Leu Met Pro Thr Pro Lys Ala Gly Thr

Val Thr Thr Asp Val Val Lys Thr Val Ala Glu Leu Arg Lys Gly Lys Ile Glu Phe Lys Ala Asp Arg Ala Gly Val Cys Asn Val Gly Val Ala Lys Leu Ser Phe Asp Ser Ala Gln Ile Lys Glu Asn Val Glu Ala Leu Cys Ala Ala Leu Val Lys Ala Lys Pro Ala Thr Ala Lys Gly Gln Tyr Leu Val Asn Phe Thr Ile Ser Ser Thr Met Gly Pro Gly Val Thr Val Asp Thr Arg Glu Leu Ile Ala Leu <210> 449 <211> 1252 <212> PRT <213> Chlamydia trachomatis serovar D Met Phe Lys Cys Pro Glu Arg Val Ser Ile Lys Lys Lys Glu Asp Ile Leu Asp Leu Pro Asn Leu Val Glu Val Gln Ile Lys Ser Tyr Lys Gln Phe Leu Gln Ile Gly Lys Leu Ala Glu Glu Arg Glu Asn Ile Gly Leu Glu Glu Val Phe Arg Glu Ile Phe Pro Ile Lys Ser Tyr Asn Glu Ala Thr Ile Leu Glu Tyr Leu Ser Tyr Asn Leu Gly Val Pro Lys Tyr Ser 65 70 75 80 Pro Glu Glu Cys Ile Arg Arg Gly Ile Thr Tyr Ser Val Thr Leu Lys Val Arg Phe Arg Leu Thr Asp Glu Thr Gly Ile Lys Glu Glu Glu Val Tyr Met Gly Thr Ile Pro Ile Met Thr Asp Lys Gly Thr Phe Ile Ile Asn Gly Ala Glu Arg Val Val Val Ser Gln Val His Arg Ser Pro Gly Ile Asn Phe Glu Gln Glu Lys His Ser Lys Gly Asn Val Leu Phe Ser Phe Arg Ile Ile Pro Tyr Arg Gly Ser Trp Leu Glu Ala Val Phe Asp 170

Ile	Asn	Asp	Leu 180	Ile	Tyr	Ile	His	Ile 185	Asp	Arg	Lys	ГЛS	Arg 190	Arg	Arg
Lys	Ile	Leu 195	Ala	Met	Thr	Phe	Ile 200	Arg	Ala	Leu	Glý	Tyr 205	Ser	Thr	Asp
Ala	Asp 210	Ile	Ile	Glu	Glu	Phe 215	Phe	Ser	Val	Glu	Glu 220	Arg	Ser	Leu	Arg
Leu 225	Glu	Lys	Asp	Phe	Val 230	Ala	Leu	Val	Gly	Lys 235	Val	Leu	Ala	Asp	Asn 240
Val	Val	Asp	Ala	Asp 245	Ser	Ser	Leu	Val	Tyr 250	Gly	Lys	Ala	Gly	Glu 255	Lys
Leu	Ser	Thr	Ala 260	Met	Leu	Lys	Arg	Ile 265	Leu	Asp	Ala	Gly	Val 270	Gln	Ser
Leu	Lys	Ile 275	Ala	Val	Gly	Ala	Asp 280	Glu	Asn	His	Pro	Ile 285	Ile	Lys	Met
Leu	Ala 290	Lys	Asp	Pro	Thr	Asp 295	Ser	Tyr	Glu	Ala	Ala 300	Leu	Lys	Asp	Phe
Tyr 305	Arg	Arg	.Leu	Arg	Pro 310	Gly	Glu	Pro	Ala	Thr 315	Leu	Val	Asn	Ala	Arg 320
Ser	Thr	Ile	Met	Arg 325	Leu	Phe	Phe	Asp	Ala 330	Lys	Arg	Tyr	Asn	Leu 335	Gly
Arg	Val	Gly	Arg 340	Tyr	Lys	Leu	Asn	Lys 345	Lys	Leu	Gly	Phe	Pro 350	Leu	Asp
Asp	Glu	Thr 355	Leu	Ser	Gln	Val	Thr 360	Leu	Arg	Lys	Glu	Asp 365	Val	Ile	Gly
Ala	Leu 370	Lys	Tyr	Leu	Ile	Arg 375	Leu	Arg	Met	Gly	Asp 380	Glu	Lys	Thr	Ser
Ile 385	Asp	Asp	Ile	Asp	His 390	Leu	Ala	Asn	Arg	Arg 395	Val	Arg	Ser	Val	Gly 400
Glu	Leu	Ile	Gln	Asn 405	His	Cys	Arg	Ser	Gly 410	Leu	Ala	Arg	Met	Glu 415	Lys
Ile	Val	Arg			Met			Phe 425		Phe	Ser	Ser	Asp 430	Thr	Leu
Thr	Pro	Gly 435	Lys	Ile	Ile	Ser	Ala 440	Lys	Gly	Leu	Val	Ser 445	Val	Leu	Lys
Asp	Phe 450	Phe	Ser	Arg	Ser	Gln 455	Leu	Ser	Gln	Phe	Met 460	Asp	Gln	Thr	Asn
Pro 465		Ala	Glu	Leu	Thr 470	His	Lys	Arg	Arg	Leu 475	Ser	Ala	Leu	Gly	Pro 480
Gly	Gly	Leu	Asn	Arg 485	Glu	Arg	Ala	Gly	Phe 490		Val	Arg	Asp	Val 495	His

Ala Ser His Tyr Gly Arg Ile Cys Pro Ile Glu Thr Pro Glu Gly Pro Asn Ile Gly Leu Ile Thr Ser Leu Ser Ser Phe Ala Lys Ile Asn Glu Phe Gly Phe Ile Glu Thr Pro Tyr Arg Val Val Arg Asp Gly Ile Val Thr Asp Glu Ile Glu Tyr Met Thr Ala Asp Val Glu Glu Glu Cys Val Ile Ala Gln Ala Ser Ala Glu Leu Asp Glu Tyr Asp Met Phe Lys Thr Pro Val Cys Trp Ala Arg Tyr Lys Gly Glu Ala Phe Glu Ala Asp Thr Ser Thr Val Thr His Met Asp Val Ser Pro Lys Gln Leu Val Ser Val Val Thr Gly Leu Ile Pro Phe Leu Glu His Asp Asp Ala Asn Arg Ala Leu Met Gly Ser Asn Met Gln Arg Gln Ala Val Pro Leu Leu Lys Thr Glu Ala Ala Ile Val Gly Thr Gly Leu Glu Gly Arg Ala Ala Lys Asp Ser Gly Ala Ile Ile Val Ala Gln Glu Asp Gly Val Val Glu Tyr Val Asp Ser Tyr Glu Ile Val Val Ala Lys Lys Asn Asn Pro Thr Leu Lys 680 Asp Arg Tyr Gln Leu Lys Lys Phe Leu Arg Ser Asn Ser Gly Thr Cys Ile Asn Gln Thr Pro Leu Cys Ser Val Gly Asp Val Val Thr His Gly Asp Val Leu Ala Asp Gly Pro Ala Thr Asp Lys Gly Glu Leu Ala Leu Gly Lys Asn Val Leu Val Ala Phe Met Pro Trp Tyr Gly Tyr Asn Phe Glu Asp Ala Ile Ile Ile Ser Glu Arg Leu Ile Lys Gln Asp Ala Tyr 760 Thr Ser Ile Tyr Ile Glu Glu Phe Glu Leu Thr Ala Arg Asp Thr Lys Leu Gly Lys Glu Glu Ile Thr Arg Asp Ile Pro Asn Val Ser Glu Glu Val Leu Ala Asn Leu Gly Glu Asp Gly Val Val Arg Ile Gly Ala Glu Val Lys Pro Gly Asp Ile Leu Val Gly Lys Ile Thr Pro Lys Ser Glu

	820			825					830		
Thr Glu Leu 835	Ala Pro	Glu Glu	Arg 840	Leu	Leu	Arg	Ala	Ile 845	Phe	Gly	Glu
Lys Ala Ala 850	Asp Val	Lys Asp 855	Ala	Ser	Leu	Thr	Val 860	Pro	Pro	Gly	Thr
Glu Gly Val 865	Val Met	Asp Val 870	Lys	Val	Phe	Ser 875	Arg	Lys	Asp	Arg	Leu 880
Ser Lys Ser	Asp Asp 885	Glu Leu	Val	Glu	Glu 890	Ala	Val	His	Leu	Lys 895	Asp
Leu Gln Lys	Glu Tyr 900	Lys Ser	Gln	Leu 905	Ala	Gln	Leu	Lys	Val 910	Glu	His
Arg Glu Lys 915	Leu Gly	Ala Leu	Leu 920	Leu	Asn	Glu	Lys	Ala 925	Pro	Ala	Ala
Ile Ile His 930	Arg Arg	Ser Ala 935	Asp	Ile	Leu	Val	Gln 940	Glu	Gly	Ala	Ile
Phe Asp Gln 945	Glu Thr	Ile Glu 950	Leu	Leu	Glu	Arg 955	Glu	Ser	Leu	Val	Asp 960
Leu Leu Met	Ala Pro 965	Cys Asp	Met	Tyr	Asp 970	Val	Leu	Lys	Asp	Ile 975	Leu
Ser Ser Tyr	Glu Thr 980	Ala Val	Gln	Arg 985	Leu	Glu	Val	Asn	Tyr 990	Lys	Thr
Glu Ala Glu 995		Lys Glu	Gly 1000		Ala	Asp	Leu	Asp 100		Gly	Val
Ile Arg Gln 1010	Val Lys	Val Tyr 101		Ala	Ser	Lys	Arg 1020		Leu	Gln	Val
Gly Asp Lys 1025	Met Ala	Gly Arg 1030	His	Gly	Asn	Lys 1035		Val	Val	Ser	Lys 1040
Ile Val Pro	Glu Ala 1049	_	Pro	Phe	Leu 1050		Asn	Gly	Glu	Thr 105	
Gln Met Ile	Leu Asn 1060	Pro Leu	Gly	Val 106		Ser	Arg	Met	Asn 1070		Gly
Gln Val Leu 107		His Leu	Gly 1080		Ala	Ala	Lys	Thr 108	_	Gly	Ile
Tyr Val Lys 1090	Thr Pro	Val Phe 109		Gly	Phe	Pro	Glu 110		Arg	Ile	Trp
Asp Met Met 1105	Ile Glu	Gln Gly 1110	Leu	Pro	Glu	Asp 1115		Lys	Ser	Tyr	Leu 1120
Phe Asp Gly	Lys Thr	_	Arg	Phe	Asp 1130		Lys	Val	Val	Val 113	_
Tyr Ile Tyr	Met Leu 1140	Lys Leu	Ser	His 114		Ile	Ala	Asp	Lys 115		His

263

Ala Arg Ser Ile Gly Pro Tyr Ser Leu Val Thr Gln Gln Pro Leu Gly 1155 1160 1165

Gly Lys Ala Gln Met Gly Gly Gln Arg Phe Gly Glu Met Glu Val Trp 1170 1175 1180

Ala Leu Glu Ala Tyr Gly Val Ala His Met Leu Gln Glu Ile Leu Thr 1185 1190 1195 1200

Val Lys Ser Asp Asp Val Ser Gly Arg Thr Arg Ile Tyr Glu Ser Ile 1205 1210 1215

Val Lys Gly Glu Asn Leu Leu Arg Ser Gly Thr Pro Glu Ser Phe Asn 1220 1225 1230

Val Leu Ile Lys Glu Met Gln Gly Leu Gly Leu Asp Val Arg Pro Met 1235 1240 1245

Val Val Asp Ala 1250

<210> 450

<211> 298

<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 450

Met Leu Lys Ile Asp Leu Thr Gly Lys Ile Ala Phe Ile Ala Gly Ile
5 10

Gly Asp Asp Asn Gly Tyr Gly Trp Gly Ile Ala Lys Met Leu Ala Glu 20 25 30

Ala Gly Ala Thr Ile Leu Val Gly Thr Trp Val Pro Ile Tyr Lys Ile 35 40 45

Phe Ser Gln Ser Leu Glu Leu Gly Lys Phe Asn Ala Ser Arg Glu Leu 50 55 60

Ser Asn Gly Glu Leu Leu Thr Phe Ala Lys Ile Tyr Pro Met Asp Ala 65 70 75 80

Ser Phe Asp Thr Pro Glu Asp Ile Pro Gln Glu Ile Leu Glu Asn Lys 85 90 95

Arg Tyr Lys Asp Leu Ser Gly Tyr Thr Val Ser Glu Val Val Glu Gln
100 105 110

Val Lys Lys His Phe Gly His Ile Asp Ile Leu Val His Ser Leu Ala 115 120 125

Asn Ser Pro Glu Ile Ala Lys Pro Leu Leu Asp Thr Ser Arg Lys Gly 130 135 140

Tyr Leu Ala Ala Leu Ser Thr Ser Ser Tyr Ser Phe Ile Ser Leu Leu 145 150 155 160

Ser His Phe Gly Pro Ile Met Asn Ala Gly Ala Ser Thr Ile Ser Leu 165 170 175

Thr Tyr Leu Ala Ser Met Arg Ala Val Pro Gly Tyr Gly Gly Met Asn Ala Ala Lys Ala Ala Leu Glu Ser Asp Thr Lys Val Leu Ala Trp Glu Ala Gly Arg Arg Trp Gly Val Arg Val Asn Thr Ile Ser Ala Gly Pro Leu Ala Ser Arg Ala Gly Lys Ala Ile Gly Phe Ile Glu Arg Met Val Asp Tyr Tyr Gln Asp Trp Ala Pro Leu Pro Ser Pro Met Glu Ala Glu Gln Val Gly Ala Ala Ala Ala Phe Leu Val Ser Pro Leu Ala Ser Ala Ile Thr Gly Glu Thr Leu Tyr Val Asp His Gly Ala Asn Val Met Gly Ile Gly Pro Glu Met Phe Pro Lys Asp <210> 451 <211> 298 <212> PRT <213> Chlamydia trachomatis serovar D Met Ser Leu Gln Lys Leu Leu Val Thr Asp Ile Asp Gly Thr Ile Thr His Gln Ser His Leu Leu His Asp Arg Val Val Lys Ala Leu His Gln Tyr Tyr Asp Ser Gly Trp Gln Leu Phe Phe Leu Thr Gly Arg Tyr Phe Ser Tyr Ala Tyr Pro Leu Phe Gln Asn Phe Ser Val Pro Phe Leu Leu Gly Ser Gln Asn Gly Ser Ser Val Trp Ser Ser Thr Asp Lys Glu Phe Ile Tyr Phe Arg Ser Leu Ser Arg Asp Phe Leu Tyr Val Leu Glu Lys Tyr Phe Glu Asp Leu Asp Leu Ile Ala Cys Ile Glu Ser Gly Ala Ser Asn Arg Asp Val Tyr Phe Arg Lys Gly Leu Gly Lys Thr Ser Gln Glu Leu Lys Ala Ile Leu Asp Ala Val Tyr Phe Pro Thr Pro Glu Ala Ala Arg Leu Leu Val Asp Val Gln Gly His Leu Ser Glu Glu Phe Ser Tyr

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Ile Leu Ser Val Asp Lys Glu Lys Gly Arg Val Ala Leu Gly Leu Lys 275 280 285

Gln Lys Glu His Asn Pro Trp Glu Asp Ile Glu Lys Lys Tyr Pro Pro 290 . 295 300

Gly Lys Arg Val Arg Gly Lys Ile Val Lys Leu Leu Pro Tyr Gly Ala 305 310 315 320

Phe Ile Glu Ile Glu Gly Ile Glu Gly Leu Ile His Val Ser Glu 325 330 335

Met Ser Trp Val Lys Asn Ile Val Asp Pro Asn Glu Val Val Asn Lys 340 345 350

Gly Asp Glu Val Glu Val Val Leu Ser Ile Gln Lys Asp Glu Gly 355 360 365

Lys Ile Ser Leu Gly Leu Lys Gln Thr Lys His Asn Pro Trp Asp Asn 370 375 380

Ile Glu Glu Lys Tyr Pro Ile Gly Leu Arg Val Thr Ala Glu Ile Lys 385 390 395 400

Asn Leu Thr Asn Tyr Gly Ala Phe Val Glu Leu Glu Pro Gly Ile Glu 405 415

Gly Leu Ile His Ile Ser Asp Met Ser Trp Ile Lys Lys Val Ser His 420 425 430

Pro Ser Glu Leu Phe Lys Lys Gly Asn Thr Val Glu Ala Val Ile Leu 435 440 445

Ser Val Asp Lys Glu Ser Lys Lys Ile Thr Leu Gly Val Lys Gln Leu 450 460

Thr Pro Asn Pro Trp Asp Glu Ile Glu Val Met Phe Pro Val Gly Ser 470 475 480

Asp Ile Ser Gly Val Val Thr Lys Ile Thr Ala Phe Gly Ala Phe Val 485 490 495

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Glu Lys Pro Phe Ala Lys Ile Glu Asp Val Leu Ser Ile Gly Asp Lys 515 520 525

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35 40 45

Glu Ile Pro Leu Ser Ser Thr Asp His Arg Thr Gly Ala Ile Trp His $50 \hspace{1cm} 55 \hspace{1cm} 60$

Ile Glu Ile Ala Gly Ile Ser Ser Glu Trp Ser Tyr Ala Tyr Lys Leu 65 70 75 80

Arg Gly Thr Asp Leu Ser Ser Gln Lys Phe Ala Thr Asp Ser Tyr Ile 85 90 95

Ala Asp Pro Tyr Ser Lys Asn Ile Tyr Ser Pro Gln Leu Phe Gly Ser 100 105 110

Pro Lys Gln Glu Lys Asp Tyr Ala Phe Ser Tyr Leu Lys His Glu Asp 115 120 125

Phe Asp Trp Glu Gly Asp Thr Pro Leu His Leu Pro Lys Glu Asn Tyr 130 135 140

Phe Ile Tyr Glu Met His Val Arg Ser Phe Thr Arg Asp Pro Ser Ser 145 150 155 160

Gln Val Ser His Pro Gly Thr Phe Leu Gly Ile Ile Glu Lys Ile Asp 165 170 175

His Leu Lys Gln Leu Gly Val His Ala Val Glu Leu Leu Pro Ile Phe 180 185 190

Glu Phe Asp Glu Thr Val His Pro Phe Lys Asn Gln Asp Phe Pro His 195 200 205

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Arg Arg Tyr Thr Tyr Gly Ala Asp Pro Cys Ala Pro Ala Arg Glu Phe 225 230 235 240

Lys Thr Leu Val Lys Ala Leu His Arg Ala Gly Ile Glu Val Ile Leu 245 250 255

Asp Val Val Phe Asn His Thr Gly Phe Glu Gly Thr Ser Cys Pro Leu 260 265 270

Pro Trp Ile Asp Leu Glu Ser Tyr Tyr Met Val Asn Asp His Gly Asp 275 280 285 .

Leu Met Asn Phe Ser Gly Cys Gly Asn Thr Val Asn Thr Asn Thr Pro 290 300

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Asn	Glu 610	Arg	Ile	Glu	Ile	Ser 615	Leu	Pro	Lys	Pro	Arg 620	Lys	Glu	His	Leu
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270

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PCT/US01/23121 WO 02/08267

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287

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Ala Gln Met Asp Gly Ala Ile Leu Val Val Ser Ala Thr Asp Gly Ala 100 $105\ 
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288

PCT/US01/23121

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Asp Thr Phe Glu Lys Leu Gly Phe Ala Leu Asp Phe Phe Ser Arg Thr
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Thr Asn Pro Phe His Ala Glu Leu Val Gln Asp Phe Tyr Ser Gln Leu
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                                                 110
Lys Ala Ser Gly Leu Ile Glu Asn Arg Ile Ser Glu Gln Leu Tyr Ser
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120

125

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Pro Arg Cys Gly Phe Asp His Ala Arg Gly Asp Glu Cys Gln Ser Cys
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Gly Ala Asp Tyr Glu Ala Ile Asp Leu Ile Gly Pro Lys Ser Lys Ile
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Ser Gly Val Glu Leu Val Lys Lys Glu Thr Glu His Ser Tyr Phe Leu
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Leu Asp Arg Met Lys Asp Ala Leu Leu Ser Phe Ile Gln Gly Cys Tyr
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Leu Pro Asp His Val Arg Lys Phe Val Val Asp Tyr Ile Glu His Val 210 215 220
Arg Ser Arg Ala Ile Thr Arg Asp Leu Ser Trp Gly Ile Pro Val Pro 225 230 235 240
Asp Phe Pro Gly Lys Val Phe Tyr Val Trp Phe Asp Ala Pro Ile Gly 245 250 255
Tyr Ile Ser Gly Thr Met Glu Trp Ala Ala Ser Gln Gly Asn Pro Asp
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Glu Trp Lys Arg Phe Trp Leu Glu Asp Gly Val Glu Tyr Val Gln Phe 275 280 285
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Glu Leu Gly Gln Lys Leu Asp Tyr Lys Lys Val Asp Ala Leu Val Val
305 310 315
Ser Glu Phe Tyr Leu Leu Glu Gly Arg Gln Phe Ser Lys Ser Glu Gly 325 330 335
Asn Tyr Val Asp Met Asp Lys Phe Leu Ser Ser Tyr Ser Leu Asp Lys 340 345 350
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Cys Tyr Arg Glu Tyr Ser Leu Arg Lys Ala Thr Ser Val Ile Met Ser
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Leu Ala Ala Leu Gly Asn Val Tyr Phe Asn Gln Gln Ala Pro Trp Lys
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<211> 97

<212> PRT

<213> Chlamydia pneumoniae

<400> 493

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Leu Val Leu Leu Asp Arg Glu Gly Ile Gln Pro Glu Phe Thr Glu Glu
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Glu Thr Ile Ser Val Thr Thr Arg Lys Pro Arg Glu Gly Glu Val Pro
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Gly Lys Asp Tyr His Phe Val Ser His Glu Glu Phe Gln Arg Leu Leu
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Asp Arg Gln Ala Leu Leu Glu Trp Val Phe Leu Phe Gly Glu Cys Tyr
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Gly Thr Ser Met Leu Glu Ile Glu Arg Ile Trp Ser Leu Gly Lys His
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Ala Val Ala Val Ile Asp Ile Gln Gly Ala Leu Phe Ile Arg Ser Arg
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Glu Arg Arg Leu Ala Ser Arg Gly Ser Glu Glu Gly Ser Gln Arg Lys
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Glu Arg Leu Glu His Ser Leu Ile Glu Leu Ala Ala Ala Asn Gln Phe
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Asp Tyr Val Ile Ile Asn Asp Asp Leu Asn Gln Ala Tyr Arg Val Leu
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Met Asp Leu Glu Arg Glu Arg Gly Ile Thr Ile Lys Ala His Pro Val
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	50					55					60				
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Gln	Ala	Gln 115	Ser	Leu	Ala	Asn	Val 120	Tyr	Leu	Ala	Leu	Glu 125	Arg	Asp	Leu
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	450					455		_	Val		460				
465					470	_	_		Ser	475	_	-	_		480
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	_	515	_			_	520	-	Leu		_	525			
Gln	Leu 530	Phe	Lys	Ile	Pro	11e 535		Ala	Ala	Ile	Asn 540	Lys	Lys	Val	Ile

PCT/US01/23121 WO 02/08267

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                                         45
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Val Gly Gly Phe Val Lys Ile Gln Gly Gln Arg Phe Val Leu Ile Gly
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 130
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145 150 155 160
Ala Tyr Pro Gly Leu Thr Ala Glu Glu Arg Gly Gln Gly Trp Ala Ile
165 170 175
Ala Lys Asn Leu Phe Glu Leu Ser Arg Leu Ala Thr Pro Val Ile Ile 180 $180\ 
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Pro Glu Gly Cys Ala Ser Ile Leu Trp Lys Asp Pro Lys Lys Asn Ser 225 230 235 240
Glu Ala Ala Ser Met Leu Lys Met His Gly Glu Asn Leu Lys Gln Phe
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Gly Ile Ile Asp Thr Val Ile Lys Glu Pro Ile Gly Gly Ala His His 260 265 270
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<212> PRT

293

<213> Chlamydia pneumoniae

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Glu Tyr Asn Lys Gly Ser Leu Arg Asn His Ile Ala Cys Val Leu Gln 470 475 Asn Pro Phe Leu Phe Tyr Asp Thr Val Trp Asn Asn Leu Thr Cys Gly 490 485 Lys Asp Met Glu Glu Glu Ala Val Leu Glu Ala Leu Lys Arg Ala Tyr 500 505 505 Ala Asp Glu Phe Ile Leu Lys Leu Pro Lys Gly Val His Ser Val Leu 515 520 525 Glu Glu Ser Gly Lys Asn Leu Ser Gly Gly Gln Gln Gln Arg Leu Ala 535 Ile Ala Arg Ala Leu Leu Lys Asn Ala Ser Ile Leu Ile Leu Asp Glu 550 555 Ala Thr Ser Ala Leu Asp Ala Ile Ser Glu Asn Tyr Ile Lys Asn Ile 565 570 Ile Gly Glu Leu Lys Gly Gln Cys Thr Gln Ile Ile Ile Ala His Lys 580 585 Leu Thr Thr Leu Glu His Val Asp Arg Val Leu Tyr Ile Glu Asn Gly 595 600 605 Gln Lys Ile Ala Glu Gly Thr Lys Glu Glu Leu Leu Gln Thr Cys Pro 615 620 Glu Phe Leu Lys Met Trp Glu Leu Ser Gly Thr Lys Glu Tyr Asn Arg 630 635 640 Val Phe Val Pro Asp His Lys Leu Val Ala Asn Pro Thr Asp Met Ala 645 650 Ile Thr Thr <210> 498 <211> 411 <212> PRT <213> Chlamydia pneumoniae <400> 498 Met Ile Pro Thr Met Leu Met Phe Phe Ile Ile Cys Phe Thr Leu Cys 10

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```
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Leu Gln Ser Ser Asp Asp Leu Leu Pro Leu Leu Lys Lys Pro Tyr Tyr 260 265 270
Met Pro Glu Thr Ile Ser Ala Lys Met Ala Leu Cys Gln Met Ala Ala
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Glu Asp Glu Thr Leu Gly Met Ile Ile Asp Glu Tyr Gly Ser Ile Glu
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                                      300
Gly Leu Ile Thr Gln Glu Asp Leu Phe Glu Ile Val Ala Gly Glu Ile
305 310
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Val Asp Gln Arg Asp Asn Lys Ile Leu Tyr Thr Thr Ser Gly Ala Asp
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                                330
Val Ile Ile Ala Ser Gly Thr Leu Glu Leu Arg Glu Phe Ser Glu Ile
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Phe Asp Ile Asn Leu Pro Thr Asn Asn Asn Ile Ala Thr Ile Gly Gly 355 360 365
Trp Leu Ile Glu Gln Ile Gly Thr Ile Pro Thr Thr Gly Met Lys Leu 370 375 380
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Arg Asn Cys Tyr Arg Ala Leu Gly Ile Thr Pro Asp Tyr Ala Pro Phe
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Thr Gln Ile Phe Ile Val Val Ile Phe Ala Glu Leu Leu Pro Leu Thr
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                                         110
Ile Ser Arg Lys Ile Pro Glu Lys Leu Ala Leu Trp Gly Ala Pro Ile
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                        120
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Leu Tyr Tyr Ser His Tyr Ile Phe Tyr Pro Leu Ile Gln Leu Ile Gly 130 140
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Ser Leu Thr Glu Gly Leu Tyr Tyr Leu Leu Asn Ile Arg Lys Glu Lys 145 150 155 160
Leu Asn Ser Thr Leu Ser Arg Asp Glu Phe Gln Lys Ala Leu Glu Thr
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                                                    175
His His Glu Glu Gln Asp Phe Asn Thr Ile Ala Thr Asn Ile Phe Ser
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                            185
Leu Ser Ala Thr Cys Ala Asp Gln Val Cys Gln Pro Leu Glu Gln Val
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      195
                                            205
Thr Met Leu Pro Ser Ser Ala Asn Val Lys Asp Phe Cys Arg Thr Ile
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                                       220
Lys Asn Thr Asp Ile Asn Phe Ile Pro Val Tyr His Lys Ala Arg Lys
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PCT/US01/23121 WO 02/08267 296

Asn Val Ile Gly Ile Ala His Pro Lys Asp Phe Val Asn Lys Ala Leu 245 250 Asp Glu Pro Leu Ile Asn Asn Leu His Ser Pro Trp Phe Ile Thr Ala 265 Lys Ser Lys Leu Ile Arg Ile Leu Lys Glu Phe Arg Asp Asn Arg Ser 285 275 280 Ser Val Ala Val Val Leu Asn Ala Ser Gly Glu Pro Ile Gly Ile Leu 295 Ser Leu Asn Ala Ile Phe Lys Ile Leu Phe Asn Thr Thr Asn Ile Ala 310 315 His Leu Lys Pro Lys Thr Ile Ser Val Ile Glu Arg Thr Phe Pro Gly 325 330 335 Asn Ser Arg Ile Lys Asp Leu Gln Lys Glu Leu Asp Ile Gln Phe Pro 345 Gln Tyr Pro Val Glu Thr Leu Ala Gln Leu Val Leu Gln Leu Leu Asp 355 360 365 360 Ser Pro Ala Glu Val Gly Thr Ser Val Ile Ile Asn Asn Leu Leu Leu 370 375 380 Glu Val Lys Glu Met Ser Leu Ser Gly Ile Lys Thr Val Ser Ile Lys 390 395 Asn Leu Leu Ser <210> 500 <211> 543 <212> PRT <213> Chlamydia pneumoniae

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Ala Trp Ser His Gln Leu Gln Asn His Lys Glu Phe Leu Ser Lys Val 265 260 Arg His Val Met Cys Thr Ser Pro Phe Ala Lys Val Arg Phe Gln Ala 280 Ala Ala Leu Leu His Leu His Gly Asp Pro Leu Gly Arg Asp Ser Leu 295 300 Val Glu Gly Leu Arg Ser Pro Gln Pro Leu Val Cys Glu Ala Ala Ser 315 310 Ala Ala Leu Cys Ser Leu Gly Ile His Gly Val Pro Leu Ala Lys Glu 325 330 His Leu Glu Ser Leu Ser Ser Arg Lys Ala Ala Ala Asn Leu Ser Ile 345 350 340 Leu Leu Leu Val Ser Arg Glu Asp Ile Glu Arg Ala Gly Asp Val Ile 355 360 365 Ala Arg Tyr Leu Ser Asn Pro Glu Met Cys Trp Ala Ile Glu Tyr Phe 375 380 Leu Trp Asp Ala Gln Trp Asn Leu Arg Gly Asp Thr Phe Pro Leu Tyr 390 395 Ser Asp Met Ile Lys Arg Glu Ile Gly Arg Lys Leu Ile Arg Leu Leu 410 Ala Val Ala Arg Tyr Ser Gln Ala Lys Ala Val Thr Ala Thr Phe Leu 425 430 420 Ser Gly Gln Gln Ala Gln Gly Trp Ser Phe Phe Ser Gly Met Phe Trp $435 \hspace{1.5cm} 440 \hspace{1.5cm} 445$ Glu Glu Gly Asp Val Lys Thr Ser Glu Asp Leu Val Thr Asp Ala Cys 455 460 Phe Ala Ala Lys Leu Glu Gly Ala Leu Ala Ser Leu Cys Gln Lys Lys 470 475 Asp Gln Ala Ser Leu Gln Arg Val Ser Gln Leu Tyr Asn Asp Ser Arg 485 490 Trp Gln Asp Lys Leu Ala Ile Leu Glu Ser Val Ala Phe Ser Glu Asn 505 510 500 Leu Asp Ala Val Pro Phe Leu Leu Asp Cys Cys His His Glu Ala Pro 520 525 Ser Leu Arg Ser Ala Ala Ala Gly Ala Leu Phe Ser Ile Phe Lys 530 <210> 501 <211> 103 <212> PRT <213> Chlamydia pneumoniae <400> 501 Met Ser Phe Lys Arg Phe Leu Gln Gln Ile Pro Val Arg Ile Cys Leu 10 Leu Ile Ile Tyr Leu Tyr Gln Trp Leu Ile Ser Pro Leu Leu Gly Ser 25 Cys Cys Arg Phe Phe Pro Ser Cys Ser His Tyr Ala Glu Gln Ala Leu 35 40 4.5 Lys Ser His Gly Phe Leu Met Gly Cys Trp Leu Ser Ile Lys Arg Ile 55 60 Gly Lys Cys Gly Pro Trp His Pro Gly Gly Ile Asp Met Val Pro Lys 70 75 Thr Ala Leu Gln Glu Val Leu Glu Pro Tyr Gln Glu Ile Asp Gly Gly 85 Asp Ser Ser His Phe Ser Glu 100 <210> 502

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<212> PRT

298

<213> Chlamydia pneumoniae

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Phe Arg Lys Asp Ile Tyr Lys Leu His Leu Phe Ser Gly Pro Leu Ile
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Ala Lys Ser Ser Arg Lys Val Tyr Leu Ser Glu Asp Phe Leu Asn Glu
Ile Ser Gln Ala Ser Leu Asp Asp Leu Ile Ser Leu Phe Lys Asp Glu
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Arg Tyr Met Tyr Gly Arg Pro Ile Lys Leu Trp Ala Leu Ser Val Ala
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Ile Ala Ser His His Ile Asp Ile Thr Pro Val Leu Ser Lys Pro Leu
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Thr Tyr Thr Glu Leu Lys Gly Ser Ser Val Arg Trp Leu Leu Pro Asn
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Ile Asp Leu Lys Asp Phe Pro Val Ile Leu Asp Tyr Leu Arg Cys His
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Lys Tyr Pro Tyr Thr Ser Lys Gly Leu Phe Leu Leu Ile Glu Lys Met
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Val Gln Glu Gly Trp Val Asp Glu Asp Cys Leu Tyr His Phe Cys Ser
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Thr Pro Glu Phe Leu Tyr Leu Arg Thr Leu Leu Val Gly Ala Asp Val
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Gln Ala Ser Ser Val Ala Ser Leu Ala Arg Met Val Ile Arg Cys Gly
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Ile Ser Ala Thr Gln Arg Gln Lys Val Leu Lys Ser Tyr Leu Asp Cys
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Glu Glu Ser Leu Ala Ala Leu Leu Leu Leu Val His Asp Ser Asp Val
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Val Leu His Glu Phe Cys Asp Glu Asp Leu Glu Lys Val Ile Arg Leu
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Met Pro Gln Glu Ser Pro Tyr Ser Gln Asn Phe Phe Ser Arg Leu Gln
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His Ser Pro Arg Arg Glu Leu Ala Cys Met Ser Thr Gln Arg Val Glu
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Ala Pro Arg Val Gln Glu Asp Gln Asp Glu Glu Tyr Val Val Gln Asp
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Gly Asp Ser Leu Trp Leu Ile Ala Lys Arg Phe Gly Ile Pro Met Asp
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PCT/US01/23121 WO 02/08267 299

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Gly	Ser	Phe	Phe 100	Ser	Leu	Gly	Ser	Leu 105	Trp	Ala	Ala	Thr	Phe 110		Cys
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Arg	Ile 130	Ala	Glu	Val	Arg	Ser 135	Arg	Phe	Phe	Leu	Glu 140	Ala	Leu	Pro	Ala
Lys 145		Met	Ala	Leu	Asp 150	Ser	Asp	Leu	Val	Ser 155	Gly	Arg	Ala	Ser	Tyr 160
			Lys	165					170				_	175	
Phe	Ser	Ala	Met 180	Glu	Gly	Val	Phe	Arg 185	Phe	Val	Lys	Gly	Asp 190	Ala	Ile
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_	210		Ser			215					220				
225	=		Leu		230					235				_	240
			Leu	245					250					255	
			Glu 260					265					270		
		275	Ile				280					285		_	
	290		Leu			295					300		_	_	
305			Ser		310					315				_	320
			Cys	325					330					335	
_			Ser 340					345			_		350		
		355	Ser Val				360					365			
_	370		Asn			375					380				
385					390					395					400
			Glu Lys	405					410					415	
			420 Arg	-				425					430		_
		435	Ala		-		440		-			445	-		
	450		Val			455				-	460				
465			Lys		470					475					480
			Ile	485					490			-		495	
			500 Ile					505					510		
OIG	-1011	515	110	9	9	- 4,2	520			J. U	Jiu	525	Det	4 CL _	

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Lys Asp Phe Arg Ala Ile Val Thr Ser Cys Glu Thr Arg Phe Glu Met
                535
Lys Lys Met Leu Asp Pro His Phe Pro Asp Leu Leu Val Leu Ser His
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Gly Cys Ser Ser Tyr Ser Ser Lys His Lys Gln Ser Leu Ile Ile Pro
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Ile His Asp Asp Pro Val Ala Phe Ser Pro Glu Gln Ala Lys Arg Ala 35 40 45
Met Asp Leu Ser Ile Ala Gln Leu Leu Phe Asp Gly Leu Thr Arg Glu
                       55
                                          60
Thr His Arg Glu Ser Asn Asp Leu Glu Leu Ala Ile Ala Ser Arg Tyr
                                       75
Thr Val Ser Glu Asp Phe Cys Ser Tyr Thr Phe Phe Ile Lys Asp Ser
           85
                                  90
Ala Leu Trp Ser Asp Gly Thr Pro Ile Thr Ser Glu Asp Ile Arg Asn 100 105 110
Ala Trp Glu Tyr Ala Gln Glu Asn Ser Pro His Ile Gln Ile Phe Gln 115 120 125
Gly Leu Asn Phe Ser Thr Pro Ser Ser Asn Ala Ile Thr Ile His Leu
                     135
Asp Ser Pro Asn Pro Asp Phe Pro Lys Leu Leu Ala Phe Pro Ala Phe
                150 155
Ala Ile Phe Lys Pro Glu Asn Pro Lys Leu Phe Ser Gly Pro Tyr Thr 165 170 175
Leu Val Glu Tyr Phe Pro Gly His Asn Ile His Leu Lys Lys Asn Pro 180 185 190
Asn Tyr Tyr Asp Tyr His Cys Val Ser Ile Asn Ser Ile Lys Leu Leu
195 200 205
Ile Ile Pro Asp Ile Tyr Thr Ala Ile His Leu Leu Asn Arg Gly Lys
                      215
                                           220
Val Asp Trp Val Gly Gln Pro Trp His Gln Gly Ile Pro Trp Glu Leu
                   230
                                       235
His Lys Gln Ser Gln Tyr His Tyr Tyr Thr Tyr Pro Val Glu Gly Ala
                                                      255
               245
                                   250
Phe Trp Leu Cys Leu Asn Thr Lys Ser Pro His Leu Asn Asp Leu Gln 260 265 270
Asn Arg His Arg Leu Ala Thr Cys Ile Asp Lys Arg Ser Ile Ile Glu
275 280 285
Glu Ala Leu Gln Gly Thr Gln Gln Pro Ala Glu Thr Leu Ser Arg Gly 290 295 300
Ala Pro Gln Pro Asn Gln Tyr Lys Lys Gln Lys Pro Leu Thr Pro Gln
                                      315
                 310
Glu Lys Leu Val Leu Thr Tyr Pro Ser Asp Ile Leu Arg Cys Gln Arg
             325
                                 330
Ile Ala Glu Ile Leu Lys Glu Gln Trp Lys Ala Ala Gly Ile Asp Leu 340 345 350
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Ile Leu Glu Gly Leu Glu Tyr His Leu Phe Val Asn Lys Arg Lys Val
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Gln Asp Tyr Ala Ile Ala Thr Gln Thr Gly Val Ala Tyr Tyr Pro Gly
 370
                      375
Ala Asn Leu Ile Ser Glu Glu Asp Lys Leu Leu Gln Asn Phe Glu Ile
               390
                                      395
Ile Pro Ile Tyr Tyr Leu Ser Tyr Asp Tyr Leu Thr Gln Asp Phe Ile
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Glu Gly Val Ile Tyr Asn Ala Ser Gly Ala Val Asp Leu Lys Tyr Thr
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Tyr Phe Pro
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Glu Glu Ser Asp Leu Gly Lys Lys Glu Thr Glu Glu Leu Glu Ala Met
     35
                          40
Lys Gln Gln Phe Val Lys Asn Ala Glu Lys Ile Glu Glu Glu Leu Thr
                      55
                                        60
Ser Ile Tyr Asn Lys Leu Gln Asp Glu Asp Tyr Met Glu Ser Leu Ser
                                  75
                70
Asp Ser Ala Ser Glu Glu Leu Arg Lys Lys Phe Glu Asp Leu Ser Gly 85 90 95
Glu Tyr Asn Ala Tyr Gln Ser Gln Tyr Tyr Gln Ser Ile Asn Gln Ser 100 \, 105 \, 110 \,
Asn Val Lys Arg Ile Gln Lys Leu Ile Gln Glu Val Lys Ile Ala Ala
                         120
                                           125
Glu Ser Val Arg Ser Lys Glu Lys Leu Glu Ala Ile Leu Asn Glu Glu
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Asp Ile Ser Gln Ala Gln Pro His His Ile Ala Phe Leu Asp Asn Glu
                         40
                                           45
Lys Tyr Ser Ser Phe Leu Lys Asn Thr Lys Ala Gly Ala Ile Ile Leu
                     55
                                      60
Ser Arg Ser Gln Ala Met Gln His Ala His Leu Lys Lys Asn Phe Leu
                70
                                     75
Ile Thr Asn Glu Ser Pro Ser Leu Thr Phe Gln Lys Cys Ile Glu Leu
                                  90
Phe Ile Glu Pro Val Thr Ser Gly Phe Pro Gly Ile His Pro Thr Ala
Val Ile His Pro Thr Ala Arg Ile Glu Lys Asn Val Thr Ile Glu Pro
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120
                                               125
Tyr Val Val Ile Ser Gln His Ala His Ile Gly Ser Asp Thr Tyr Ile
                135
                                           140
Gly Ala Gly Ser Val Ile Gly Ala His Ser Val Leu Gly Ala Asn Cys
          150
                             155
Leu Ile His Pro Lys Val Val Ile Arg Glu Arg Val Leu Met Gly Asn
165 170 175
Arg Val Val Val Gln Pro Gly Ala Val Leu Gly Ser Cys Gly Phe Gly
                                        190
                           185
Tyr Ile Thr Asn Ala Phe Gly His His Lys Pro Leu Lys His Leu Gly
                         200
Tyr Val Ile Val Gly Asp Asp Val Glu Ile Gly Ala Asn Thr Thr Ile
                      215
                                          220
Asp Arg Gly Arg Phe Lys Asn Thr Val Ile His Glu Gly Thr Lys Ile 225 230 230 235
Asp Asn Gln Val Gln Val Ala His His Val Glu Ile Gly Lys His Ser
245 250 255
Ile Ile Val Ala Gln Ala Gly Ile Ala Gly Ser·Thr Lys Ile Gly Glu
260 265 270
                               265
His Val Ile Ile Gly Gly Gln Thr Gly Ile Thr Gly His Ile Ser Ile
                          280
Ala Asp His Val Ile Met Ile Ala Gln Thr Gly Val Thr Lys Ser Ile
                    295
                                          300
Thr Ser Pro Gly Ile Tyr Gly Gly Ala Pro Ala Arg Pro Tyr Gln Glu
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                                       315
Thr His Arg Leu Ile Ala Lys Ile Arg Asn Leu Pro Lys Thr Glu Glu 325 330 335
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Leu Ala Glu Ile Pro Ser Glu Ile
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                           40
Gln Asp Leu Thr Asn Pro Ala Ala Ala Thr Arg Thr Lys Lys Glu
Glu Lys Phe Gln Thr Leu Glu Ser Arg Lys Lys Gly Glu Ala Gly Lys 65 70 75 80
Ala Glu Lys Lys Ser Glu Ser Thr Glu Glu Lys Pro Asp Thr Asp Leu
85 90 95
Ala Asp Lys Tyr Ala Ser Gly Asn Ser Glu Ile Ser Gly Gln Glu Leu
          100
                               105
Arg Gly Leu Arg Asp Ala Ile Gly Asp Asp Ala Ser Pro Glu Asp Ile
                          120
Leu Ala Leu Val Gln Glu Lys Ile Lys Asp Pro Ala Leu Gln Ser Thr
                       135
                                 140
Ala Leu Asp Tyr Leu Val Gln Thr Thr Pro Pro Ser Gln Gly Lys Leu
                150
                             155
Lys Glu Ala Leu Ile Gln Ala Arg Asn Thr His Thr Glu Gln Phe Gly
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                         170
Arg Thr Ala Ile Gly Ala Lys Asn Ile Leu Phe Ala Ser Gln Glu Tyr
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Ala Asp Gln Leu Asn Val Ser Pro Ser Gly Leu Arg Ser Leu Tyr Leu
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Glu Val Thr Gly Asp Thr His Thr Cys Asp Gln Leu Leu Ser Met Leu
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                                        220
Gln Asp Arg Tyr Thr Tyr Gln Asp Met Ala Ile Val Ser Ser Phe Leu
               230 - 235
Met Lys Gly Met Ala Thr Glu Leu Lys Arg Gln Gly Pro Tyr Val Pro
245 250 255
Ser Ala Gln Leu Gln Val Leu Met Thr Glu Thr Arg Asn Leu Gln Ala
                             265
Val Leu Thr Ser Tyr Asp Tyr Phe Glu Ser Arg Val Pro Ile Leu Leu
                         280
Asp Ser Leu Lys Ala Glu Gly Ile Gln Thr Pro Ser Asp Leu Asn Phe
                           300
 290 295
Val Lys Val Ala Glu Ser Tyr His Lys Ile Ile Asn Asp Lys Phe Pro
     310
                          315
Thr Ala Ser Lys Val Glu Arg Glu Val Arg Asn Leu Ile Gly Asp Asp 325 330 335
Val Asp Ser Val Thr Gly Val Leu Asn Leu Phe Phe Ser Ala Leu Arg
                            345
Gln Thr Ser Ser Arg Leu Phe Ser Ser Ala Asp Lys Arg Gln Gln Leu
     355
                         360
                                            365
Gly Ala Met Ile Ala Asn Ala Leu Asp Ala Val Asn Ile Asn Asn Glu
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Lys Gln Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr Leu His Ile Lys
                         40
Ala Phe Pro Cys Tyr Leu Ser Ala Leu His Ala Arg Cys Asp Asp Leu
                     55
                                        60
Gln Ile Arg Arg Gln Ile Leu Glu Asn Leu Met Asp Glu Glu Ala Gly
                  70
                                     75
Asn Pro Asn His Ile Asp Leu Trp Arg Gln Phe Ala Leu Ser Leu Gly
                                 90
Val Ser Glu Glu Glu Leu Ala Asn His Glu Phe Ser Gln Ala Ala Gln
                            105
Asp Met Val Ala Thr Phe Arg Arg Leu Cys Asp Met Pro Gln Leu Ala 115 120 125
Val Gly Leu Gly Ala Leu Tyr Thr Tyr Glu Ile Gln Ile Pro Gln Val
                   135
 130
Cys Val Glu Lys Ile Arg Gly Leu Lys Glu Tyr Phe Gly Val Ser Ala
                 150
                                    155
Arg Gly Tyr Ala Tyr Phe Thr Val His Gln Glu Ala Asp Ile Lys His
               165
                                170
                                                   175
Ala Ser Glu Glu Lys Glu Met Leu Gln Thr Leu Val Gly Arg Glu Asn
                             185
        180
Pro Asp Ala Val Leu Gln Gly Ser Gln Glu Val Leu Asp Thr Leu Trp
                         200
Asn Phe Leu Ser Ser Phe Ile Asn Ser Thr Glu Pro Cys Ser Cys Lys
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210
                        215
                                            220
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 Ile Val Val Ile Thr Ser Lys Ile Val Ser Leu Cys Glu Gly Ala Val
                            40
 Val Glu Leu Glu Lys Val Ser Lys Asp Glu Leu Ile Lys Gln Glu Ala
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                                           60
 Asp Ala Tyr Val Phe Val Glu Lys Tyr Gly Ile Tyr Leu Thr Lys Lys
                  70
                                       75
 Trp Gly Ile Leu Ile Pro Ser Ala Gly Ile Asp Glu Ser Asn Val Glu
                85
                                   90
 Gly Tyr Phe Val Leu Tyr Pro Arg Asp Phe Leu Leu Ser Val Asn Thr
                                105
 Leu Gly Asp Trp Leu Arg Asn Phe Tyr His Leu Glu His Cys Gly Ile
                            120
                                               125
 Ile Ile Ser Asp Ser His Thr Thr Pro Leu Arg Arg Gly Thr Met Gly
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                                            140
 Leu Gly Leu Cys Trp Asn Gly Phe Phe Pro Leu Tyr Asn Tyr Val Gly
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                                    155
 Lys Pro Asp Cys Phe Gly Arg Ala Leu Lys Met Thr Tyr Ser Asn Leu 165 170 175
 Leu Asp Gly Leu Ser Ala Ala Ala Val Leu Cys Met Gly Glu Gly Asp
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                                185
                                                   190
 Glu Gln Thr Pro Ile Ala Ile Ile Glu Glu Ala Pro Lys Ile Thr Phe
                            200
 His Ser Ser Pro Thr Thr Leu Gln Asp Met Ser Thr Leu Ala Ile Ala
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                                           220
 Glu Asp Glu Asp Leu Tyr Gly Pro Leu Leu Gln Ser Met Ala Trp Glu
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 Thr Pro Ala Pro Thr Ser
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                                                    30
 Phe Lys Thr Phe Leu Lys Ser Glu Glu Ala Ile Ile Tyr Ser Asn Gln
                             40
                                                45
 Cys Asn Glu Asp Met Arg Lys Ile Leu Cys Asp Ala Ile Glu His Ala
                         55
 Asp Glu Glu Ile Phe Leu Arg Ile Tyr Asn Leu Ser Glu Pro Lys Ile
                     70
                                        75
 Gln Gln Ser Leu Thr Arg Gln Ala Gln Ala Lys Asn Lys Val Thr Ile
                                    90
 Tyr Tyr Gln Lys Phe Lys Ile Pro Gln Ile Leu Lys Gln Ala Ser Asn
                                105
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PCT/US01/23121 WO 02/08267

305

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Val Thr Leu Val Glu Gln Pro Pro Ala Gly Arg Lys Leu Met His Gln
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      115
Lys Ala Leu Ser Ile Asp Lys Lys Asp Ala Trp Leu Gly Ser Ala Asn
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Tyr Thr Asn Leu Ser Leu Arg Leu Asp Asn Asn Leu Ile Leu Gly Met
              150
                                  155
His Ser Ser Glu Leu Cys Asp Leu Ile Ile Thr Asn Thr Ser Gly Asp
                            170
           165
Phe Ser Ile Lys Asp Gln Thr Gly Lys Tyr Phe Val Leu Pro Gln Asp
                           185
         180
                                          190
Arg Lys Ile Ala Ile Gln Ala Val Leu Glu Lys Ile Gln Thr Ala Gln
                       200
                                        205
     195
Lys Thr Ile Gln Val Ala Met Phe Ala Leu Thr His Ser Glu Ile Ile
                                   220
                   215
Gln Ala Leu His Gln Ala Lys Gln Arg Gly Ile His Val Asp Ile Ile
    230
                                235
Ile Asp Arg Ser His Ser Lys Leu Thr Phe Lys Gln Leu Arg Gln Leu
                             250
           245
Asn Ile Asn Lys Asp Phe Val Ser Ile Asn Thr Ala Pro Cys Thr Leu
                           265 270
His His Lys Phe Ala Val Ile Asp Asn Lys Thr Leu Leu Ala Gly Ser
       275
                     280
Ile Asn Trp Ser Lys Gly Arg Phe Ser Leu Asn Asp Glu Ser Leu Ile
290 295 300
Ile Leu Glu Asn Leu Thr Lys Gln Gln Asn Gln Lys Leu Arg Met Ile
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                       315
Trp Lys Asp Leu Ala Lys His Ser Glu His Pro Thr Val Asp Asp Glu
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Glu Lys Glu Ile Ile Glu Lys Ser Leu Pro Val Glu Glu Gln Glu Ala
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<211> 186
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<213> Chlamydia pneumoniae
<400> 511
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Arg Leu Ile Gly Ser Ala Gly Glu Gln Leu Gly Ile Leu Ala Ile Lys
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Asp Ala Leu Asp Leu Ala Arg Glu Ala Gly Leu Asp Leu Val Glu Val
      35
                        40
Ala Ser Asn Ser Glu Pro Pro Val Cys Lys Ile Met Asp Tyr Gly Lys
                   55
                                    60
Tyr Arg Tyr Gly Leu Thr Lys Lys Glu Lys Asp Ser Lys Lys Ala Gln
                                   75
                 70
His Gln Val Arg Ile Lys Glu Val Lys Leu Lys Pro Asn Ile Asp Glu
85 90 95
                               90
Asn Asp Phe Ser Thr Lys Leu Lys Gln Ala Arg Thr Phe Val Glu Lys
          100
                           105 110
Gly Asn Lys Val Lys Ile Thr Cys Met Phe Arg Gly Arg Glu Leu Ala
    115
                        120
                                          125
Tyr Pro Glu His Gly Phe Lys Val Val Gln Lys Met Ser Gln Gly Leu
 130 135
                                140
Glu Asp Ile Gly Phe Val Glu Ala Glu Pro Lys Leu Ala Gly Arg Ser 145 150 155
Leu Ile Cys Val Val Ala Pro Gly Thr Val Lys Thr Lys Lys Gln
                                                 175
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Glu Lys Ser His Ala Gln Asp Glu Asn Gln
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Glu Val Ile Ile Gly Thr Asp Thr Thr Pro Thr Val Thr Lys Phe Ser
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Gly Asp Lys Gly Ile Val Ile Thr Thr Asp Ser Thr Ile Thr Pro Ser
                   55
                                     60
Ser Thr Thr Phe Ser Leu Asp Met Glu Ala Val Ile Lys Glu Val Thr
                70
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Asp Lys Ile Leu Thr Gln Ile Glu Asp Glu Leu Val Lys Asp Ile Ile
              85
                                90
Lys Asn Ile Thr Gln Ser Leu Ile Glu Glu Val Ile Lys Lys Ile His
       100 105
                                             110
Ile Asp Pro Ser Phe Ser Tyr Ser Arg Ala Phe Lys Asp Val Asn Ile
                                     125
                        120
Thr Asn Lys Ile Gln Cys Asn Gly Leu Phe Thr Lys Glu Asn Ile Gly
                    135
                                      140
Asn Leu Asp Gly Gly Thr Glu Ile Ala Ser Ser Ser Val Thr Pro Asp
                  150
                                    155
Asn Ala Asn Ser Met Phe Leu Ile Cys Ala Asp Ile Ile Ala Thr Arg
              165
                                170
                                                175
Met Glu Gly Thr Val Ala Leu Ala Leu Val Lys Glu Gly Asp Leu Ser
    180
                          185
                                       190
Pro Cys Ser Ile Ser Tyr Gly Tyr Ser Ala Gly Tyr Pro Asn Ile Ile
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                                         205
Ser Leu Arg Ala Thr Val Gly Asn Lys Thr Thr Ala Pro Val Lys Phe
                    215
                                        220
Ser Leu Arg Ala Gly Gly Met Asp Ser Gly Val Val Trp Val Asn Ala
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                                    235
Met Pro Asn Gly Glu Lys Ile Leu Gly Val Asp Ala Val Ser Lys Ile
245 250 255
Thr Ile Leu Glu Val Lys Pro Gln Thr Asn Gly Thr Xaa Xaa Xaa Xaa
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Phe Xaa Xaa Xaa
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<400> 513
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Asp	Leu 50	Glu	Glu	Ser	Ile	Ala 55	Gln	Ile	Thr	His	Lys 60	Val	Val	Гуѕ	Glu
Val 65	Leu	Ala	Lys	Ile	Ser 70	Glu	Gly	Gln	Val	Val 75	Thr	Val	Glu	Arg	Ile 80
Gln	Asp	Leu	Val	Glu 85	Ser	Gln	Leu	Tyr	Ile 90	Ser	Gly	Leu	Gln	Asp 95	Val
Ala	Arg	Asp	Tyr 100	Ile	Val	Tyr	Arg	Asp 105	Gln	Arg	Lys	Ala	Glu 110	Arg	Gly
Asn	Ser	Ser 115	Ser	Ile	Ile	Ala	Ile 120	Ile	Arg	Arg	Asp	Gly 125	Gly	Ser	Ala
Lys	Phe 130	Asn	Pro	Met	Lys	Ile 135	Ser	Ala	Ala	Leu	Glu 140	Lys	Ala	Phe	Arg
145	Thr				150	_				155					160
	Asn			165					170					175	
~	Glu		180					185			_		190		-
	Leu	195					200					205			
_	Arg 210					215					220				
225	Glu				230					235					240
-	Gly			245				_	250	_				255	
	Trp		260	_	_			265			_		270		
	Asp	275		•			280					285			
	Thr 290					295					300				
305	Asp	Tyr	Ата	rne	310	Ата	Ата	GIU	ьеп	315	Thr	ser	ser	ren	320
Glu	Glu	Thr	Leu	Gly 325	Суз	Ser	Ser	Gln	Asp 330	Pro	Asn	Leu	Ser	Glu 335	Ile
	Lys		340					345					350		
	Asn	355					360					365			
	Asp 370					375					380				
385	Tyr	_			390					395	_	-			400
	Gln			405					410					415	
_	Glu		420					425		•	-		430		
	Phe	435					440					445		_	
_	His 450					455					460				
465	Ser			_	4.70				_	475					480
_	Ala	_		485					490			_		495	
	Val		500				Gly	505	ı		_		510		
Ile	Lys	val	Ala	Asn	Asp	rnr	АТа	тте	ATa	Val	Asn	Gln	GLy	GTÀ	туѕ

		515					520					525			
Arg	Lys 530	Gly	Ala	Met	Cys	Val 535		Leu	Glu	Asn	Trp 540		Leu	Asp	Tyr
Glu 545		Phe	Leu	Glu	Leu 550	Arg	Lys	Asn	Thr	Gly 555	Asp	Glu	Arg	Arg	Arg 560
Thr				565					570					Phe 575	
Arg	Leu	Glu	Lys 580	Lys	Gly	Met	Trp	Thr 585	Leu	Phe	Ser	Pro	Asp 590	Asp	Val
Pro	Gly	Leu 595	His	Glu	Ala	Tyr	Gly 600	Leu	Glu	Phe	Glu	Lys 605	Leu	Tyr	Glu
	610		_			615					620			Lys	
625					630					635				Tyr	640
	_			645					650					Arg 655	
			660					665					670	Thr	
		675					680					685		Leu	
	690					695					700			Glu	
705		-			710					715				Asn	720
	_			725					730					Asn 735	
		-	740					745					750	Val	
-		755					760					765		Phe	
	770					775					780			Ser	
785			_		790					795				Ser	800
_	_			805					810					Lys 815	
	_	_	820					825					830	Lys	
		835					840					845		Asn	
	850					855					860			Ile	
865					870					875				Lys	880
				885					890					Lys 895	
	_		900					905					910		Lys
-		915					920					925			Leu
_	930					935					940				Ile
945	_				950					955					Leu 960
				965					970					Met 975	
			980)				985					990		
GID	нта	995		ser	val	GIU	100		FIIE	116	nsp	100			Arg

309

Gly Ile Gln Pro Arg Trp Met Lys Asn Lys Ser Ala Ser Thr Ser Ile 1020 1010 1015 Val Val Glu Arg Lys Thr Thr Pro Val Cys Ser Met Glu Glu Gly Cys 1030 Glu Ser Cys Gln <210> 514 <211> 346 <212> PRT <213> Chlamydia pneumoniae <400> 514 Met Glu Ala Asp Ile Leu Asp Gly Lys Leu Lys Arg Val Glu Val Ser 10 Lys Lys Gly Leu Val Asn Cys Asn Gln Val Asp Val Asn Gln Leu Val Pro Ile Lys Tyr Lys Trp Ala Trp Glu His Tyr Leu Asn Gly Cys Ala 35 40 45Asn Asn Trp Leu Pro Thr Glu Val Pro Met Ala Arg Asp Ile Glu Leu 55 Trp Lys Ser Asp Glu Leu Ser Glu Asp Glu Arg Arg Val Ile Leu Leu 70 Asn Leu Gly Phe Phe Ser Thr Ala Glu Ser Leu Val Gly Asn Asn Ile 85 90 Val Leu Ala Ile Phe Lys His Ile Thr Asn Pro Glu Ala Arg Gln Tyr 100 105 110 Leu Leu Arg Gln Ala Phe Glu Glu Ala Val His Thr His Thr Phe Leu 115 120 Tyr Ile Cys Glu Ser Leu Gly Leu Asp Glu Gly Glu Val Phe Asn Ala 130 135 140 135 140 Tyr Asn Glu Arg Ala Ser Ile Arg Ala Lys Asp Asp Phe Gln Met Thr 150 155 Leu Thr Val Asp Val Leu Asp Pro Asn Phe Ser Val Gln Ser Ser Glu 165 170 Gly Leu Gly Gln Phe Ile Lys Asn Leu Val Gly Tyr Tyr Ile Ile Met 180 185 190 Glu Gly Ile Phe Phe Tyr Ser Gly Phe Val Met Ile Leu Ser Phe His 195 200 205 Arg Gln Asn Lys Met Thr Gly Ile Gly Glu Gln Tyr Gln Tyr Ile Leu 210 215 220 Arg Asp Glu Thr Ile His Leu Asn Phe Gly Ile Asp Leu Ile Asn Gly 225 230 235 240 Ile Lys Glu Glu Asn Pro Glu Val Trp Thr Thr Glu Leu Gln Glu Glu 245 250 Ile Val Ala Leu Ile Glu Lys Ala Val Glu Leu Glu Ile Glu Tyr Ala 260 265 Lys Asp Cys Leu Pro Arg Gly Ile Leu Gly Leu Arg Ser Ser Met Phe 280 285 Ile Asp Tyr Val Arg His Ile Ala Asp Arg Arg Leu Glu Arg Ile Gly 290 295 300 Leu Lys Pro Ile Tyr His Ser Arg Asn Pro Phe Pro Trp Met Ser Glu 310 315 Thr Met Asp Leu Asn Lys Glu Lys Asn Phe Phe Glu Thr Arg Val Thr 325 330 Glu Tyr Gln Thr Ala Gly Asn Leu Ser Trp . 340 345 <210> 515 <211> 327

<212> PRT

310

<213> Chlamydia pneumoniae

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<212> PRT

<213> Chlamydia pneumoniae

<400> 516

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 Phe Ser Lys Thr Glu
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 Thr Leu Ser Leu Asn Val Gly Cys Asn Leu Ala Gln Tyr Ser Asn Lys 20 25 30
 Lys Val Leu Leu Val Asp Leu Asp Pro Gln Ala Asn Leu Thr Thr Gly
                           40
 Leu Gly Val Gln Ser Cys Tyr Glu Ser Asn Leu Asn Asp Ile Phe Arg 50 55 60
 Ser Ser Gly Asn Val Arg Asp Ile Ile Gln Asp Thr Lys Ile Glu Asn
                                        75
 Leu His Ile Val Pro Ser Ser Ile Leu Ile Glu Glu Phe Arg Glu Phe
                                     90
              85
 Asn Arg Asn Ser Val Leu Asp Thr Ser His Leu Arg Ser Ser Leu Gln 100 105 110
 Leu Ile Glu Ser Asn Tyr Asp Leu Cys Ile Leu Asp Thr Pro Pro Ser 115 120 125
                                        125
 Leu Gly Thr Leu Thr Glu Glu Ala Phe Ile Ala Ser Asp His Leu Ile
130 135
                                           140
 Val Cys Leu Thr Pro Glu Pro Phe Ser Ile Leu Gly Leu Gln Lys Ile
145 150 155 160
 Lys Glu Phe Cys Ser Val Leu Pro Lys Lys Lys Asp Leu Ser Val Leu
165 170 175
 Gly Ile Val Phe Ser Phe Trp Asp Gly Arg Asn Ser Thr Asn Ser Thr
            180
                                  185
                                                      190
 Tyr Leu Asn Ile Ile Glu Ser Ile Tyr Glu Gly Lys Val Leu Ser Ser
                                                205
                          200
 Lys Val Arg Arg Asp Ile Thr Leu Ser Arg Ser Leu Leu Lys Glu Thr 210 215 220
                                             220
 Ser Ile Ala Asn Ala Tyr Pro Asn Ser Arg Ala Ser His Asp Ile Leu
225 230 235 240
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 Arg Leu Thr Lys Glu Ile Glu Asp Lys Leu Phe Asn Lys Glu Met Ser
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 Ala Gln Glu Val Leu
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 <213> Chlamydia pneumoniae
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Lys Leu Ile Gly Thr Ser Pro Lys His Gly Ile Tyr Leu Pro Leu Phe 20 25
 Ser Ile His Thr Lys Asn Ser Cys Gly Ile Gly Glu Phe Leu Asp Leu 35 40 45
 Ile Pro Leu Ile Ser Trp Cys Gln Lys Gln Gly Phe Ser Val Ile Gln
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Leu 65	Leu	Pro	Leu	Asn	Asp 70	Thr	Gly	Glu	Asp	Thr 75	Ser	Pro	Tyr	Asn	Ser
Ile	Ser	Ser	Val	Ala 85	Leu	Asn	Pro	Leu	Phe 90	Leu	Ser	Leu	Ser	Ser 95	
Pro	Asn	Ile	Asp 100		Ile	Pro	Glu	Val 105		Lys	Lys	Leu	Gln 110		Met
His	Glu	Leu 115	Cys	Ser	Thr	Pro	Ser 120		Ser	Tyr	Thr	Gln 125		Lys	Glu
Lys	Lys 130		Ala	Phe	Leu	Arg 135		Tyr	Tyr	Gln	Lys 140		Суз	Lys	Ser
Ser 145	Leu	Glu	Gly	Asn	Ser 150		Phe	Ser	Glu	Phe 155		Glu	Ser	Glu	Arg 160
Tyr	Trp	Leu	Tyr	Pro 165	Tyr	Gly	Thr	Phe	Arg 170		Ile	Lys	His	His 175	
His	Gly	Glu	Pro 180	Ile	Asn	Asn	Trp	Pro 185		Ser	Leu	Thr	Asp 190		Glu
Asn	Phe	Pro 195	Asp	Leu	Thr	Lys	Lys 200	Phe	His	Asp	Glu	Val 205		Phe	Phe
	210		Gln			215					220				
225			Gln		230					235					240
			Asp	245					250					255	
			Ser 260					265					270		
		275	His				280					285			
	290		Trp			295					300				_
305			Arg		310					315					320
			Ser	325					330					335	
			Gln 340					345					350		
		355	Pro				360					365		_	
	370		Leu			375					380				
385			Asn		390					395					400
			Leu	405					410			_		415	
			Trp 420					425				-	430		
		435	His				440					445			
	450		Leu			455					460				
465			Asn		470					475					480
			Arg	485					490					495	Lys
			Tyr 500					505					510	Ile	His
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Asp Leu Ala Val Ser Thr Leu Leu Gly Thr Leu Pro Glu Asn Val Phe
                    55
Arg Glu Arg Ile Phe Lys Ala Ala Leu Ser Val Asn Gly Ser Phe Gln 65 70 75 80
Ser Ser Ile Lys Gly Ile Leu Gly Tyr Gly Glu Val Thr Gln Gln Leu
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Tyr Leu Ser Asp Ile Leu Ser Met Asn Tyr Leu Asn Gly Glu Lys Leu 100 105 110
Phe Glu Tyr Leu Lys Leu Phe Ser Leu His Ala Lys Ile Trp Met Glu 115 120 125
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Tyr 'Val Ala
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Leu Asp Gln Tyr Arg Thr Ile Val Ser Ala Ile Ser Thr Ala Leu Lys
                       40
Glu Asn Ala Ala Phe Lys Ala Asn Thr Leu Thr Gln Ile Val Pro Leu
               55
                                         60
Asn Val Asp Val Leu Ser Leu Phe Ser Asp Val Leu Asp Leu Asp Ala
                   70
                                      75
Gly Ile Pro Glu Thr Pro Asn Val Leu Leu Ser Asn Glu Met Gln Lys
                                90
Val Phe Gln Gly Ile Tyr Asn Glu Ile Ser Leu Ile Lys Val Phe Pro
                             105
                                                110
Asn Gly Asp Lys Ile Val Val Ala Ser Ser Ile Pro Glu His Leu Gly
       115
                        120
                                             125
Glu Asn Tyr Asn His Lys Ile Asp Ile Pro Lys Asn Thr Pro Phe Leu
                     135
Ala Ala Leu Lys Gln Ser Pro Lys Asn Gln Glu Val Phe Ser Val Met
                  150
                                     155
Gln Ala Asn Val Phe Asp Ala Lys Thr Gln Glu Leu Gln Gly Ile Leu
                               170
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Tyr Thr Thr Phe Ser Ala Glu Ser Leu Leu Lys Asp Leu Leu Ile Asn
          180
                             185
Lys Gln Ser Tyr Leu Thr Val Lys Thr Ala Ile Leu Ser Lys Tyr Gly
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                                          205
Val Ile Leu Lys Ala Ser Asp Pro Ala Leu His Leu His Thr Val Tyr
 210 . 215
                                         220
Pro Asp Met Thr Lys Glu Lys Phe Cys Gln Val Phe Leu Asn Asp Asp
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Pro	Cys	Pro	Ile	Asp 245	Ser	Glu	Leu	Gly	Pro 250	Leu	Thr	Leu	Ser	Pro 255	Leu
Asp	Ile	Gly	Glu 260		Phe	Tyr	Ser	Phe 265		Ile	Lys	Asp	Thr 270		Ile
Trp	Gly		Ile	Glu	Asn	Val			Ile	Asp	Ile		-	Leu	Ser
Tyr		275 Lys	Lys	Glu	Glu		280 Phe	Ala	Pro	Leu		285 Arg	Arg	Ala	Arg
Met	290 Tyr	Thr	Ala	Tyr		295 Phe	Cys	Ile	Leu		300 Gly	Ser	Leu	Ile	
305 Phe	Ile	Val	Ala	Arg	310 Arg	Leu	Ser	Leu	Pro	315 Ile	Arg	Lys	Leu	Ala	320 Thr
			Glu	325					330					335	
			340					345					350		
		355	Phe				360					365			
Val	Glu 370	Asn	Leu	His	Lys	Gln 375	Gln	His	Leu	Ala	Lys 380	Thr	Asn	Phe	Glu
Met 385	Lys	Glu	Asn	Ala	Gln 390	Asn	Ala	Leu	His	Leu 395	Gly	Glu	Gln	Ala	Gln 400
	Arg	Leu	Leu	Pro 405	Asn	Thr	Leu	Pro	Ser 410		Pro	His	Ile	Glu 415	Leu
Ala	Lys	Ala	Tyr 420		Pro	Ala	Ile	Thr 425		Gly	Gly	Asp	Phe 430	-	Asp
Val	Phe	Val 435	Val	Gly	Glu	Gly	Ser 440		Ala	Arg	Leu	Phe 445		Ile	Val
Ala	Asp 450		Ser	Gly	Lys	Gly 455		Asn	Ala	Суѕ	Gly 460		Ser	Leu	Phe
Leu 465	Lys	Asn	Met	Leu	Arg 470	Thr	Phe	Leu	Ser	Arg 475	Ser	Ser	Ser	Leu	Gln 480
	Ala	Ile	Gln	Glu 485		Ser	Arg	Leu	Phe 490	-	Asn	Asn	Thr	Lys 495	
Ser	Gly	Met	Phe 500		Thr	Leu	Cys	Val 505		Cys	Tyr	His	Gln 510		Ser
Asn	Thr	Met 515	Glu	Tyr	Tyr	Ser	Cys 520		His	Pro	Pro	Ala 525		Tyr	Leu
Asp	Pro 530		Gly	Glu	Thr	Ser 535		Leu	Phe	His	Pro 540		Met	Ala	Leu
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Ile	Gln		580 Leu	Thr	Gly	Lys		585 Ala	Ala	Asp	Ala		590 His	Arg	Leu
Met	Leu	595 Ser	Val	Lys	Thr	Phe	600 Val	Gly	Asn	Ser	His	605 Gln	His	Asp	Asp
Ile	610 Thr	Leu	Leu	Ile	Leu	615 Lys	Val	Leu	Glu	Ser	620				
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Pro Ile Ser Ala Ile Leu Gly Leu Gln Ser Leu Phe Leu Ser Ile Gly
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Gly Gly Ile Ala Leu Gly Thr Ile Ala Ala Leu Lys Lys Lys Gln
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Arg Arg Tyr Ile Leu Gly Ala Ser Ile Leu Gln Ile Ser Ile Pro Ala
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Leu Leu Pro Ile Ala Cys Trp Gly Ser Phe Thr His Thr Ile Leu Pro
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Thr Leu Ala Leu Ala Val Thr Pro Met Ala Phe Ile Ile Gln Leu Thr
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Tyr Ser Ser Val Ser Ala Ala Leu Asn Lys Asp Tyr Val Leu Leu Ala
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Tyr Ala Lys Gly Leu Ser Pro Leu Lys Val Val Ile Lys His Ile Leu
                                       220
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Pro Tyr Ala Ile Phe Pro Thr Ile Ser Tyr Ser Ala Phe Leu Thr Thr
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       230
Thr Val Ile Thr Gly Thr Phe Ala Ile Glu Asn Ile Phe Cys Ile Pro
                   250 255
           245
Gly Leu Gly Lys Trp Phe Ile Cys Ser Ile Lys Gln Arg Asp Tyr Pro
260 265 270
Val Ala Leu Gly Leu Ser Val Phe Tyr Gly Thr Leu Phe Met Leu Ser
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Ser Leu Leu Ser Asp Leu Ile Gln Ser Ile Ile Asp Pro Gln Ile Arg
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Asp Phe Tyr Lys Glu Cys Thr Gln Lys Gly Ile Gln Pro Ile Ile Gly
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Cys Glu Cys Tyr Ile Ala Pro Gly Ser Arg Phe Asp Lys Lys Glu
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Lys Arg Ser Arg Ala Ala His His Leu Ile Leu Cys Lys Asn Glu
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Gln Gly Tyr Arg Asn Leu Cys Ile Leu Thr Ser Leu Ala Phe Thr Glu
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                            105
                                              110
Gly Phe Tyr Tyr Phe Pro Arg Ile Asp Lys Asp Leu Leu Arg Gln Tyr
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Ser Glu Gly Leu Ile Cys Leu Ser Gly Cys Leu Ser Ser Ser Val Ser
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Lys	Met	Ser	Glu 180	165 Glu	Ser	Ile	Ala	Gly 185	170 Phe	Lys	Glu	Glu	Trp 190		Lys
Gln	Glu	Tyr 195	Tyr	Ser	Leu	Ile	Glu 200		Gln	Ile	Lys	Val 205		Thr	Ala
Val	Leu 210		Ala	Ser	Lys	Arg 215		Gly	Ile	Pro	Thr 220		Ala	Thr	Asn
Asp 225		His	Tyr	Ile	Asn 230		Asn	Asp	Trp	Gln 235		His	Glu	Ile	Leu 240
	Asn	Val	Gln	Ser 245		Glu	Thr	Val	Arg 250		Ala	Lys	Gln	Asn 255	
His	Ile	Pro	Asn 260		Lys	Arg	Lys	Val 265		Arg	Ser	Arg	Glu 270		Tyr
Phe	Lys	Ser 275	Pro	Ala	Gln	Met	Ala 280		Leu	.Phe	Lys	Asp 285		Pro	Glu
Val	Ile 290	Ser	Asn	Thr	Leu	Glu 295	Val	Ala	Lys	Arg	Cys 300	Asp	Phe	Thr	Phe
Asp 305	Phe	Ser	Lys	Lys	His 310	Tyr	Pro	Ile	Tyr	Val 315	Pro	Glu	Ser	Leu	Lys 320
Thr	Leu	Asn	Ser	Tyr 325	Thr	Glu	Glu	Asp	Arg 330	Tyr	Gln	Ala	Ser	Ala 335	Val
Phe	Leu	Lys	Gln 340	Leu	Ala	Glu	Glu	Ala 345	Leu	Pro	Lys	Lys	Tyr 350	Ser	Ser
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Asp	Ile 370	Val	ГÀг	Glu	Arg	Met 375	Asp	Met	Glu	Met	Ala 380	Ile	Ile	Ile	Pro
Lys 385	Gly	Met	Суѕ	Asp	Tyr 390	Leu	Leu	Ile	Val	Trp 395	Asp	Ile	Ile	His	Trp 400
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_			Leu 420					425					430		
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	450		Ile			455					460				
465		-	Ala		470					475					480
			Gly	485					490					495	
			Asp 500					505					510		
		515					520					525			Pro
-	530		Gln			535					540				
545			Leu		550					555					560
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		595	Glu				600					605			
_	610		Thr			615					620				
Thr	GTÀ	GID	Ser	ьeu	ATS	тет	нта	inr	ьeu	rro	ьeu	Asp	Asp	ATS	TUL

PCT/US01/23121 **WO** 02/08267 317

625	Db	O	7	T	630	C1 =	C1	T	m}	635	G1	T1_	Dh.	C1	640
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Phe	Glu	Glu 675	Ile	Ile	Ala	Met	Gly 680	Ala	Leu	Tyr	Arg	Pro 685	Gly	Pro	Met
Asp	Met 690	Ile	Pro	Ser	Phe	Ile 695	Asn	Arg	Lys	His	Gly 700	Lys	Glu	Ile	Ile
Glu 705	Tyr	Asp	His	Pro	Leu 710	Met	Glu	Ser	Ile	Leu 715	Lys	Glu	Thr	Tyr	Gly 720
	Met	Val	Tyr	Gln 725	Glu	Gln	Val	Met	Gln 730		Ala	Gly	Ala	Leu 735	
Ser	Tyr	Ser	Leu 740	Gly	Glu	Gly	Asp	Val 745	Leu	Arg	Arg	Ala	Met 750	Gly	Lys
Lys	Asp	Phe 755	Gln	Gln	Met	Glu	Gln 760		Arg	Glu	Lys	Phe 765	Cys	Lys	Arg
Ala	Cys 770		Asn	Gly	Ile	Asp 775		Glu	Leu	Ala	Thr 780		Ile	Phe	Asp
Lys 785	Met	Glu	Lys	Phe	Ala 790	Ala	Tyr	Gly	Phe	Asn 795	Lys	Ser	His	Ala	Ala 800
Ala	Tyr	Gly	Leu	Ile 805	Thr	Tyr	Thr	Thr	Ala 810	Tyr	Leu	Lys	Ala	Asn 815	Tyr
Pro	Lys	Glu	Trp 820	Leu	Ala	Ala	Leu	Leu 825	Thr	Cys	Asp	Ser	Asp 830	Asp	Ile
Glu	Lys	Ile 835	Gly	Lys	Leu	Ile	Arg 840	Glu	Ala	Gln	Ser	Met 845	Gly	Ile	Pro
Ile	Leu 850	Pro	Pro	His	Ile	Asn 855	Val	Ser	Ser	Asn	His 860	Phe	Val	Ala	Thr
Asp 865	Glu	Gly	Ile	Arg	Phe 870	Ala	Met	Gly	Ala	Ile 875	Lys	Gly	Ile	Gly	Arg 880
				885	Ile				890					895	
			900		Phe			905					910		
Lys	Lys	Ser 915	Ile	Glu	Ser	Leu	11e 920	Asp	Ala	Gly	Суѕ	Phe 925	Asp	Суѕ	Phe
Asp	Ser 930	Asn	Arg	Asp	Leu	Leu 935	Leu	Ala	Ser	Val	Glu 940	Pro	Leu	Tyr	Glu
945			-	-	Lys 950	-				955	-				960
				965	Met				970					975	
	_	_	980		Thr			985	_				990		
		995			Ile		1000)				1005	5		
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		1075	5		Gln		1080)				1085	5		
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Arg 1105	_	Met	Lys	Asp	Leu 111(Ile	Val	Asn	Glu 1115		Ile	Ile	Tyr	Glu 1120

318

Cys Asp Gln Ala Phe Asp Arg Ile Lys Asn Gln Val Gln Lys Met Ser 1125 1130 Phe Thr Met Ser Thr Ser Gly Lys Glu Thr Lys Ala Lys Gly Asn Lys 1140 1145 1150 Pro Asn Glu Asn Gly His Thr Gln Ala Leu Ala Pro Val Thr Leu Ser 1155 1160 1165 Leu Asp Leu Asn Glu Leu Arg His Ser His Leu Cys Ile Leu Lys Lys 1170 1175 1180Ile Val Gln Lys His Pro Gly Ser Arg Thr Leu Val Leu Val Phe Thr 1185 1190 1195 1200 Gln Asp Asn Glu Arg Val Ala Ser Met Ser Pro Asp Asp Ala Tyr Phe 1205 1210 1215 Val Cys Glu Asp Ile Glu Glu Leu Arg Gln Glu Leu Val Thr Ala Asp 1220 1225 Leu Pro Val Arg Val Ile Thr Val 1235 <210> 523 <211> 576 <212> PRT <213> Chlamydia pneumoniae <400> 523 Met Thr Asp Phe Pro Thr His Phe Lys Gly Pro Lys Leu Asn Pro Ile 10 Lys Val Asn Pro Asn Phe Phe Glu Arg Asn Pro Lys Val Ala Arg Val Leu Gln Ile Thr Ala Val Val Leu Gly Ile Ile Ala Leu Leu Ser Gly 40 45 Ile Val Leu Ile Ile Gly Thr Pro Leu Gly Ala Pro Ile Ser Met Ile 55 60 Leu Gly Gly Cys Leu Leu Ala Ser Gly Gly Ala Leu Phe Val Gly Gly 70 75 Thr Ile Ala Thr Ile Leu Gln Ala Arg Asn Ser Tyr Lys Lys Ala Val Asn Gln Lys Lys Leu Ser Glu Pro Leu Met Glu Arg Pro Glu Leu Lys 100 105 Ala Leu Asp Tyr Ser Leu Asp Leu Lys Glu Val Trp Asp Leu His His 120 115 125 Ser Val Val Lys His Leu Lys Lys Leu Asp Leu Asn Leu Ser Lys Thr 135 Gln Arg Glu Val Leu Asn Gln Ile Lys Ile Asp Asp Glu Gly Pro Ser 150 155 Leu Gly Glu Cys Ala Ala Met Ile Ser Glu Asn Tyr Asp Ala Cys Leu 165 170 175 Lys Met Leu Ala Tyr Arg Glu Glu Leu Leu Lys Glu Gln Thr Gln Tyr 180 185 190 Gln Glu Thr Arg Phe Asn Gln Asn Leu Thr His Arg Asn Lys Val Leu 200 Leu Ser Ile Leu Ser Arg Ile Thr Asp Asn Ile Ser Lys Ala Gly Gly 215 220 Val Phe Ser Leu Lys Phe Ser Thr Leu Ser Ser Arg Met Ser Arg Ile 230 235 His Thr Thr Thr Val Ile Leu Ala Leu Ser Ala Val Val Ser Val 245 250 Met Val Val Ala Ala Leu Ile Pro Gly Gly Ile Leu Ala Leu Pro Ile 260 265 Leu Leu Ala Val Ala Ile Ser Ala Gly Val Ile Val Thr Gly Leu Ser 280 285 Tyr Leu Val Arg Gln Ile Leu Ser Asn Thr Lys Arg Asn Arg Gln Asp 30Ō 295

319 ·

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Tyr Ile Thr Asn Ala Pro Ile Glu Lys Arg Leu Ile Glu Glu Ile Arg
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Val Thr Tyr Lys Glu Ile Asp Ala Gln Thr Lys Lys Met Lys Thr Asp
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Leu Glu Phe Leu Glu Asn Glu Val Arg Ser Gly Arg Leu Ser Val Ala
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Ser Pro Ser Glu Asp Pro Ser Glu Thr Pro Ile Phe Thr Gln Gly Lys
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Glu Phe Ala Lys Leu Arg Arg Gln Thr Ser Gln Asn Ile Ser Thr Ile
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Tyr Gly Pro Asp Asn Glu Asn Ile Asp Pro Glu Phe Ser Leu Pro Trp
       435
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Met Pro Lys Lys Glu Glu Glu Ile Asp His Ser Leu Glu Pro Val Thr
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Lys Leu Glu Pro Gly Ser Arg Glu Glu Leu Leu Val Glu Gly Val
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Asn Pro Thr Leu Arg Glu Leu Asn Met Arg Ile Ala Leu Leu Gln Gln 485 490 495
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Gln Leu Ser Ser Val Arg Lys Trp Arg His Pro Arg Gly Glu His Tyr
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Gly Asn Val Ile Tyr Ser Asp Thr Glu Leu Asp Arg Ile Gln Met Leu
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Glu Gly Ala Phe Tyr Asn His Leu Arg Glu Ala Gln Glu Glu Ile Thr
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322

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Arg Gly Met Val Val Cys Leu Pro Asn Ser Val Lys Pro His Thr Gln
                   295 300
Phe Lys Cys Ala Val Tyr Val Leu Gln Lys Glu Glu Gly Gly Arg His
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Lys Pro Phe Phe Thr Gly Tyr Arg Pro Gln Phe Phe Phe Arg Thr Thr
              325
                               330
                                                  335
Asp Val Thr Gly Val Val Thr Leu Pro Glu Gly Ile Glu Met Val Met
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                  345
Pro Gly Asp Asn Val Glu Phe Glu Val Gln Leu Ile Ser Pro Val Ala
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Leu Glu Glu Gly Met Arg Phe Ala Ile Arg Glu Gly Gly Arg Thr Ile
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Gly Pro Leu His Phe Gly His Ile Thr Gly Ala Tyr Leu Pro Ala Asp
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Val Tyr Ala Arg Phe Gln Arg Leu Gln Gly Lys Glu Val Leu Tyr Ile
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Cys Gly Ser Asp Glu Tyr Gly Ile Ala Ile Thr Leu Asn Ala Glu Leu
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Ala Gly Met Gly Tyr Gln Glu Tyr Val Asp Met Tyr His Lys Leu His
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                                  75
Lys Asp Thr Phe Lys Lys Leu Gly Ile Ser Val Asp Phe Phe Ser Arg
85 90 95
Thr Thr Asn Thr Tyr His Pro Ala Ile Val Gln Asp Phe Tyr Arg Asn
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Leu Gln Glu Arg Gly Leu Val Glu Asn Gln Val Thr Glu Gln Leu Tyr
      115
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                                         125
Ser Glu Glu Glu Gly Lys Phe Leu Ala Asp Arg Tyr Val Val Gly Thr
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Cys Pro Lys Cys Gly Phe Asp Arg Ala Arg Gly Asp Glu Cys Gln Gln
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Cys Gly Ala Asp Tyr Glu Ala Arg Asp Leu Lys Glu Pro Arg Ser Lys
165 170 175
Leu Thr Gly Ala Ala Leu Ser Leu Arg Asp Thr Glu His Ala Tyr Leu
                          185
                                              190
His Leu Glu Arg Met Lys Glu Asp Leu Leu Ala Phe Val Gln Gly Ile
      195
                        200
Tyr Leu Arg Pro His Met Arg Asn Phe Val Thr Asp Tyr Ile Glu His
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Leu Arg Pro Arg Ala Val Thr Arg Asp Leu Ser Trp Gly Ile Pro Val
225 230 235
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340

PCT/US01/23121

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Pro Asp Leu Glu Asn Lys Val Phe Tyr Val Trp Phe Asp Ala Pro Ile
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Gly Tyr Ile Ser Gly Thr Met Asp Trp Ala Ala Ser Ile Gly Asp Pro
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Glu Ala Trp Lys Lys Phe Trp Leu Asp Asp Thr Val Thr Tyr Ala Gln
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Phe Ile Gly Lys Asp Asn Thr Ser Phe His Ala Ala Ile Phe Pro Ala
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                              300
Met Glu Ile Gly Gln Ser Leu Pro Tyr Lys Lys Val Asp Ala Leu Val
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Thr Ser Glu Phe Leu Leu Glu Gly Phe Gln Phe Ser Lys Ser Asp
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Gly Asn Phe Ile Asp Met Asp Ala Phe Leu Glu Thr Tyr Ser Leu Asp
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Lys Leu Arg Tyr Val Leu Ala Ala Ile Ala Pro Glu Thr Ser Asp Ser
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Glu Phe Ser Phe Gln Glu Phe Lys Thr Arg Cys Asn Ser Glu Leu Val
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Gly Lys Tyr Gly Asn Phe Val Asn Arg Val Leu Ala Phe Ala Val Lys
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                                 395 400
Asn Gly Cys Thr Glu Leu Ser Ser Pro Gln Leu Glu Gln Lys Asp Leu
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Asp Phe Ile Ser Lys Ser Gln Lys Leu Ala Lys Asp Ala Ala Glu His
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Tyr Ala Gln Tyr Ser Leu Arg Lys Ala Cys Ser Thr Ile Met Glu Leu
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Ala Ala Leu Gly Asn Gly Tyr Phe Asn Asp Glu Ala Pro Trp Lys Leu
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Ala Lys Glu Gly Asn Trp Asn Arg Val Arg Ala Ile Leu Phe Cys Ala 465 470 475 480
Cys Tyr Cys Gln Lys Leu Leu Ala Leu Ile Ser Tyr Pro Ile Met Pro
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Glu Thr Ala Leu Lys Ile Leu Glu Met Ile Ala Pro His Ser Leu Asp
                           505 510
Leu Gly Ser Gln Asp Pro Asp Arg Leu Gln Ser Leu Trp Thr Asp Ser
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Lys Ile Ala Arg Gly Asp Val Arg Ser Ser Asn Val Ala Ile Glu Ala
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Leu Asn Phe Leu Asp Leu Tyr Gly Ile Gln Ser Glu Tyr Ala Glu Arg
                    55
Asp Asp Arg Glu Arg His Leu Ser Ala Thr Gly Glu Arg Arg Glu
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Gln Gly Phe Gly Thr Ser Arg Arg Lys Asp Pro Ser Leu Tyr Asn Trp
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WO 02/08267

341

PCT/US01/23121

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35 40
                   40
Thr Val Ser Ser Thr Thr Arg Ser Ala Arg Pro Gly Glu Val His Gly
                      55
Val Asp Tyr Leu Phe Val Ser Glu Asp Asp Phe Lys Gln Ser Leu Asp
                 70
                                      75
Arg Glu Asp Phe Leu Glu Trp Val Phe Leu Phe Gly Thr Tyr Tyr Gly
                                90
        85
Thr Ser Lys Ala Glu Ile Ser Arg Val Leu Gln Lys Gly Lys His Cys 100 105 110
Ile Ala Val Ile Asp Val Gln Gly Ala Leu Ala Leu Lys Lys Gln Met
Pro Ala Val Thr Ile Phe Ile Gln Ala Pro Ser Gln Glu Glu Leu Glu
                    135 140
Arg Arg Leu Asn Ala Arg Asp Ser Glu Lys Asp Phe Gln Lys Lys Glu 145 150 150 160
Arg Leu Glu His Ser Ala Val Glu Ile Ala Ala Ala Ser Glu Phe Asp
165 170 175
Tyr Val Val Val Asn Asp Asp Leu Ile Thr Ala Tyr Gln Val Leu Arg
180 185 190
Ser Ile Phe Ile Ala Glu Glu His Arg Met Ser His Gly
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<213> C. Trachomatis D serovar
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His Ile Asp His Gly Lys Ser Thr Ile Ala Asp Arg Leu Leu Glu Ser
                               25
Thr Ser Thr Ile Glu Gln Arg Glu Met Arg Glu Gln Leu Leu Asp Ser
Met Asp Leu Glu Arg Glu Arg Gly Ile Thr Ile Lys Ala His Pro Val
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Thr Met Thr Tyr Glu Tyr Glu Gly Glu Thr Tyr Glu Leu Asn Leu Ile 65 70 75 80
Asp Thr Pro Gly His Val Asp Phe Ser Tyr Glu Val Ser Arg Ser Leu 85 90 95
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Gln Ala Gln Ser Leu Ala Asn Val Tyr Leu Ala Leu Glu Arg Asp Leu
115 120 125
       115
                        120
Glu Ile Ile Pro Val Leu Asn Lys Ile Asp Leu Pro Ala Ala Gln Pro
                     135
                                        140
Glu Ala Ile Lys Lys Gln Ile Glu Glu Phe Ile Gly Leu Asp Thr Ser
             150
                                    155
Asn Thr Ile Ala Cys Ser Ala Lys Thr Gly Gln Gly Ile Pro Glu Ile
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342

165 170 Leu Glu Ser Ile Ile Arg Leu Val Pro Pro Pro Lys Pro Pro Gln Glu 185 ______190 Thr Glu Leu Lys Ala Leu Ile Phe Asp Ser His Tyr Asp Pro Tyr Val 205 195 200 Gly Ile Met Val Tyr Val Arg Val Ile Ser Gly Glu Ile Lys Lys Gly 215 220 Asp Arg Ile Thr Phe Met Ala Thr Lys Gly Ser Ser Phe Glu Val Leu 230 235 Gly Ile Gly Ala Phe Leu Pro Glu Ala Thr Leu Met Glu Gly Ser Leu 245 250 Arg Ala Gly Gln Val Gly Tyr Phe Ile Ala Asn Leu Lys Lys Val Lys 260 265 270 Asp Val Lys Ile Gly Asp Thr Val Thr Thr Val Lys His Pro Ala Lys 275 280 Glu Pro Leu Glu Gly Phe Lys Glu Ile Lys Pro Val Val Phe Ala Gly 295 300 Ile Tyr Pro Ile Asp Ser Ser Asp Phe Asp Thr Leu Lys Asp Ala Leu 310 315 Gly Arg Leu Gln Leu Asn Asp Ser Ala Leu Thr Ile Glu Gln Glu Asn 325 330 335 Ser His Ser Leu Gly Phe Gly Phe Arg Cys Gly Phe Leu Gly Leu Leu 340 345 His Leu Glu Ile Ile Phe Glu Arg Ile Ser Arg Glu Phe Asp Leu Asp 360 365 Ile Ile Ala Thr Ala Pro Ser Val Ile Tyr Lys Val Val Leu Lys Asn 375 380 Gly Lys Thr Leu Phe Ile Asp Asn Pro Thr Ala Tyr Pro Asp Pro Ala 390 395 Leu Ile Glu His Met Glu Glu Pro Trp Val His Val Asn Ile Ile Thr 405 410 Pro Gln Glu Tyr Leu Ser Asn Ile Met Ser Leu Cys Met Asp Lys Arg 420 425 430 Gly Ile Cys Leu Lys Thr Asp Met Leu Asp Gln His Arg Leu Val Leu 435 440 445 Ser Tyr Glu Leu Pro Leu Asn Glu Ile Val Ser Asp Phe Asn Asp Lys 450 460 450 455 460 Leu Lys Ser Val Thr Lys Gly Tyr Gly Ser Phe Asp Tyr Arg Leu Gly 470 475 Asp Tyr Lys Lys Gly Ala Ile Ile Lys Leu Glu Ile Leu Ile Asn Asp 485 490 495 Glu Ala Val Asp Ala Phe Ser Cys Leu Val His Arg Asp Lys Ala Glu 500 505 510 Ser Lys Gly Arg Ser Ile Cys Glu Lys Leu Val Asp Val Ile Pro Pro 515 520 525 Gln Leu Phe Lys Ile Pro Ile Gln Ala Ala Ile Asn Lys Lys Ile Ile 530 535 540 540 Ala Arg Glu Thr Ile Arg Ala Leu Ala Lys Asn Val Thr Ala Lys Cys 550 555 Tyr Gly Gly Asp Ile Thr Arg Lys Arg Lys Leu Trp Asp Lys Gln Lys 565 570 Lys Gly Lys Lys Arg Met Lys Glu Phe Gly Lys Val Ser Ile Pro Asn 580 585 590 Thr Ala Phe Val Glu Val Leu Lys Met Glu

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<211> 324

<212> PRT

<213> C. Trachomatis D serovar

343

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Leu	Val 130	Val	Val	Ala	Ile	Leu 135	Lys	Ser	Thr	Thr	Leu 140		Phe	Gln	Arg
Phe 145	Leu	Ala	Gln	Leu	Ile 150	Ala	Ile	Arg	Val	Ser 155		Ser	Leu	Arg	Lys 160
Asp	Tyr	Phe	Leu	Ala 165		Gln	Thr	Leu	Pro 170		Thr	Phe	Phe	His 175	
His	Asp	Met	Gly 180	Asn	Leu	Ser	Ser	Arg 185		Ile	Ala	Asp	Ser 190		Met
Ile	Ala	Leu 195	Ala	Ile	Asn	Ala	Leu 200	Met	Val	Asn	Tyr	Ile 205		Ala	Pro
Ile	Thr 210	Met	Thr	Leu	Ala	Leu 215	Val	Val	Суѕ	Leu	Ser 220		Ser	Trp	Lys
Phe 225	Суѕ	Ala	Cys	Val	Cys 230	Leu	Ala	Phe	Pro	Ile 235	Phe	Ile	Leu	Pro	Ile 240
				245			Lys		250					255	
			260				Ala	265		_			270	_	
		275					Thr 280					285			
	290					295	Ala				300				
305					310		Leu			315					320
				325			Gly		330					335	
			340				Gly	345					350		
		355					Asn 360					365	_	_	
	370					375	Val				380				
385					390		Glu			395					400
				405			Gly		410					415	
			420				Lys	425					430		
		435					Ser 440					445			_
	450					455	Glu				460				
465					470		Leu			475			•		480
				485			Asp		490					495	
			500				Glu	505					510		
		515					Lys 520				_	525			
	530					535	Leu				540				
545					550		His			555					560
GIU	ATA	Tnr	ser	Ala 565	ьeu	Asp	Ala	Ile	Ser 570	GLu	Asn	Tyr	Val	Lys 575	Glu

345

Ile Val Gly Gln Leu Lys Gly Arg Cys Thr Gln Ile Ile Ile Ala His 580 585 Lys Leu Ser Thr Leu Glu Tyr Val Asp Arg Ile Val Tyr Leu Glu Gln 595 600 Gly Lys Lys Ile Ala Glu Gly Thr Lys Glu Glu Leu Leu Asp Ser Cys 615 620 Pro Ala Phe Gln Arg Met Trp Val Leu Ser Gly Ala Lys Asp Trp Glu 630 Leu Asn Ala Val Val Lys 645 <210> 568 <211> 414 <212> PRT <213> C. Trachomatis D serovar Met Phe Ser Ser Ala Ile Val Ile Leu Thr Ala Ile Phe Val Leu Cys 10 Ser Gly Phe Val Ser Leu Ser His Ile Ala Leu Phe Ser Leu Pro Ser 20 Ser Leu Ile Ala His Tyr Ser His Ser Lys Asn Arg Gln Leu Arg Gln 40 Ile Ala Asn Leu Met Ala Tyr Pro Asn His Leu Leu Met Thr Leu Val 55 Phe Phe Asp Ile Gly Ile Asn Ile Gly Val Gln Asn Cys Ile Ala Thr 70 Leu Val Gly Asp Ser Ala Ser Leu Leu Leu Thr Val Gly Val Pro Leu 85 90 Ala Leu Thr Leu Val Leu Gly Glu Ile Val Pro Lys Val Ile Ala Ile 100 105 Pro Tyr Asn Ala Arg Ile Ala Lys Ile Val Thr Pro Ile Ile Phe Ala 125 120 Ser Thr Lys Ser Phe Arg Pro Ile Phe Asp Trp Ala Ile Ser Gly Ile 135 140 Asn Phe Ile Val Gln Lys Met Leu Ala Arg Gln Glu Ser Asp Phe Ile 150 155 Gln Pro Gln Glu Leu Lys Glu Val Leu Arg Ser Cys Lys Asp Phe Gly 165 170 175 Val Val Asn His Glu Glu Ser Arg Leu Leu Phe Gly Tyr Leu Ser Met 180 185 Glu Glu Gly Ser Ile Lys Glu Arg Met Thr Pro Lys Gln Glu Ile Ile 195 200 205 Phe Tyr Asp Val Leu Thr Pro Ile Glu Asn Leu Tyr Lys Leu Phe Ser 215 220 Gly Pro Lys Gln Ser Tyr Ser Lys Val Leu Val Cys Lys Gly Gly Leu 230 235 Gln Asn Leu Leu Gly Val Cys Ser Ala Lys Leu Leu Leu Leu Tyr Lys 245 250 255 Glu Lys Leu Gln Ser Ala Glu Glu Leu Leu Pro Leu Leu Arg Lys Pro 260 265 His Tyr Ile Pro Glu Thr Val Ser Ala Lys Thr Ala Leu Tyr His Leu 280 Ala Gly Glu Asp Cys Gly Leu Gly Ile Ile Ile Asp Glu Tyr Gly Ser 295 300 Ile Glu Gly Leu Ile Thr Gln Asn Asp Leu Phe Lys Ile Val Ser Asp 310 315 Gly Val Ala His Asn Arg Pro Ser Phe Lys Gln Phe Ala His Ser Asp 325 330 335 Lys Asn Val Val Ile Ala Ala Gly Thr Tyr Glu Leu Ser Asp Phe Tyr 345

Asp Leu Phe Gly Val Asp Leu Pro Thr Thr Ala Asn Cys Val Thr Ile 360 Gly Gly Trp Leu Thr Glu Gln Leu Gly Glu Ile Pro Glu Thr Gly Thr 370 375 Lys Phe Ala Trp Gly Gln Phe Val Phe Gln Ile Leu Asp Ala Ala Pro 390 395 Asn Cys Val Lys Arg Val Tyr Ile Arg Lys Thr His Gly Asn . 410 <210> 569 <211> 404 <212> PRT <213> C. Trachomatis D serovar Met Glu Thr Asn Ser Pro Phe Phe Trp Leu Gly Val Asn Leu Leu Cys 10 Ile Phe Val Gln Gly Phe Phe Ser Met Met Glu Met Ala Cys Ile Ser 25 Phe Asn Arg Val Arg Leu Gln Tyr Tyr Leu Thr Lys Ser Asn Lys Lys 35 40 Ala Ser Tyr Ile Asn Phe Leu Val Arg Arg Pro Tyr Arg Leu Phe Gly 55 60 Thr Val Met Leu Gly Val Asn Ile Ala Leu Gln Ile Gly Ser Glu Ser . 70 75 Ser Arg Thr Cys Tyr Lys Leu Leu Gly Ile Ser Pro Glu Tyr Ala Pro 90 85 Ala Thr Gln Ile Ile Leu Val Val Ile Phe Ala Glu Leu Ile Pro Leu 105 Ala Ile Ser Arg Lys Ile Pro Glu Lys Ile Ala Leu Lys Gly Ala Pro 125 120 Ile Leu Tyr Phe Ala His Tyr Leu Phe Tyr Pro Leu Ile Gln Cys Val 135 Gly Gly Ile Thr Asn Met Ile Tyr Phe Ile Leu Asn Ile Lys Glu Glu 150 155 Thr Leu His Ser Thr Leu Ser Arg Asp Glu Leu Gln Lys Thr Leu Glu 165 170 Thr His His Glu Glu His Asp Phe Asn Val Ile Ala Thr Asn Ile Phe 180 185 190 Ser Leu Ser Ala Thr Ser Val Glu Gln Val Cys Gln Tyr Leu Asp Gln 195 200 205 Ile Pro Ile Leu Ser Ala Thr Ala Ser Val Arg Asp Val Cys Gln Leu 215 220 Val Arg Arg His Arg Leu Asp Phe Val Pro Val Tyr His Lys Val Lys 230 235 Lys Asn Val Val Gly Ile Ala Phe Pro Lys Asn Leu Ile Asn Arg Asn 245 250 255 Pro Ser Asp Pro Val Val Pro Tyr Leu Ser Ser Pro Trp Phe Ile Thr 265 260 Ala Lys Ser Lys Leu Ile His Ala Ile Gln Glu Phe Arg Lys Asn Ser 280 Ser Asn Val Ala Ile Val Leu Asn Asn Gly Glu Pro Met Gly Val 295 Leu Gly Leu His Thr Val Phe Lys Thr Leu Phe Asn Thr Arg Asn Ile 310 315 Ala Gln Leu Lys Pro Lys Pro Thr Ser Leu Ile Glu Arg Thr Phe Ser 325 330 Gly Asn Thr Pro Leu Ser Glu Ile Glu Asn Glu Leu Asp Ile Ile Phe 345 Met Asp Asn Asp Cys Thr Thr Ile Glu Gln Leu Met Leu Lys Leu Leu

360

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85 90 Val Ile Thr Ile Leu Ala Val Gly Met Ala Arg Asp Tyr Arg Leu Val 105 110 Pro Ile Val Leu Gln Ala Leu Ser Asp Asp Ser Asp Thr Val Arg Glu 115 120 125 Ile Ala Val Gln Val Ala Val Met Tyr Gly Ser Ser Cys Leu Leu Arg 135 Ala Val Gly Asp Leu Ala Lys Asn Asp Ser Ser Ile Gln Val Arg Ile 150 Thr Ala Tyr Arg Ala Ala Ala Val Leu Glu Ile Gln Asp Leu Val Pro 170 His Leu Arg Val Val Val Gln Asn Thr Gln Leu Asp Gly Thr Glu Arg 180 . 185 190 Arg Glu Ala Trp Arg Ser Leu Cys Val Leu Thr Arg Pro His Ser Gly 200 195 205 Val Leu Thr Gly Ile Asp Gln Ala Leu Met Thr Cys Glu Met Leu Lys 215 220 Glu Tyr Pro Glu Lys Cys Thr Glu Glu Gln Ile Arg Thr Leu Leu Ala 230 235 Ala Asp His Pro Glu Val Gln Val Ala Thr Leu Gln Ile Ile Leu Arg 245 250 Gly Gly Arg Val Phe Arg Ser Ser Ile Met Glu Ser Val Gln Lys 260 265 Leu Ala Cys Asn Ser Leu Ser Ala Arg Val Gln Met Gln Ala Ala Ala 280 Ile Leu Tyr Leu Glu Gly Asp Pro Phe Gly Glu Asp Lys Leu Thr Glu 295 300 Gly Leu Ser Ala Thr Ser Ser Ile Leu Cys Glu Ala Ala Ser Glu Ala 310 315 Val Cys Ser Leu Gly Ile His Gly Val His Leu Ala Gly Arg Phe Leu 330 325 Ser Lys Val Gln Gly Met Arg Ser Arg Val Asn Leu Ala Phe Ala Leu 345 Leu Val Ser Arg Glu Lys Val Glu Glu Ala Gly Asp Val Val Ala Ser 360 365 Phe Ile His Arg Ile Glu Pro Cys Arg Ala Ile Glu Gln Phe Leu Cys

Glu Asp Gln Lys Ile Phe Val Ala Ser Ser Pro Leu Gln Val Glu Ile 390 395 Met Lys Arg Asp Leu Ala Lys Lys Ile Ile Arg Leu Leu Val Ala Ala 405 410 Gln Tyr Ser Lys Ala Lys Met Val Val Ala Gln Tyr Leu Ala Gly Gln 420 425 430 Gln Val Gly Trp Ser Phe Cys Ser Glu Val Phe Trp Glu Glu Gly Asp 440 Ser Glu Asp Phe Val Glu Pro Leu Gln Glu Glu Ser Phe Ala Phe Ala 455 460 Leu Glu Lys Ala Leu Ser Phe Leu Gln Arg Glu Gly Gly Glu Ala Gly 470 475 Leu His Ala Val Ile Ser Leu Tyr Pro His Ser Arg Trp Gln Asp Lys 485 490 Leu Thr Ile Leu Glu Ala Ile Ala Tyr Ser Glu Asn Arg Ile Ala Thr 505 Cys Phe Leu Arg Glu Arg Cys Leu Gln Glu Ala Ala Ser Leu Gln Ser 520 Ala Ala Ala Gly Ala Val Phe Ala Leu Phe Lys 530 535 <210> 571 <211> 104 <212> PRT <213> C. Trachomatis D serovar <400> 571 Met Gln Thr Ser Arg Ile Ser Ser Phe Phe Arg Gly Leu Val His Leu 10 Tyr Arg Trp Ala Ile Ser Pro Phe Leu Gly Ala Pro Cys Arg Phe Phe 20 25 Pro Thr Cys Ser Glu Tyr Ala Leu Val Ala Leu Lys Lys His Pro Leu 40 Arg Lys Ser Leu Phe Leu Ile Ala Lys Arg Leu Leu Lys Cys Gly Pro 50 55 Trp Cys Ile Gly Gly Ile Asp Leu Val Pro Arg Thr Ser Val Glu Glu 70 75 Tyr Leu Ser Ser Pro Thr Pro Leu Ala Glu Ser Pro Asp Asp Arg Thr 85 Val Pro His Thr Gln Glu Thr Ser 100 <210> 572 <211> 336 <212> PRT <213> C. Trachomatis D serovar <400> 572 Met Gln Leu Phe Phe Gly Arg Phe Tyr Glu Val Ala Cys Ile Val Ala 5 10 Ser Ile Leu Arg Glu Arg Asp Val Gly Val Phe Met Gly Ile Glu Gly 20 25 Arg Gly Ser Gly Ala Met Gln Ser Lys Lys Thr Ile Lys Trp Leu Lys 40 Gln Ala Leu Val Leu Ser Ser Ile Val Asn Ile Leu Leu Leu Leu Leu 55 60 Ile Tyr Ser Thr Val Phe Arg Lys Asp Ile Tyr Lys Leu Arg Val Phe 70 75 Pro Gly Asn Leu Ile Ala Lys Ser Ser Arg Ile Gly Lys Ile Pro Glu 90 Asp Ile Leu Glu Arg Leu Glu Asn Ala Ser Phe Ala Asp Leu Leu Ala

105 Leu Leu Gln Glu Glu Arg Met Val Phe Gly His Pro Leu Lys Ser Trp 120 125 Ala Leu Gly Val Ser Ile Gln Lys Tyr Phe Val Asp Ile Ala Pro Met 135 140 Leu Thr His Pro Leu Thr Phe Ile Arg Leu Lys Ser Pro Glu Arg Thr 145 150 155 160 Trp Leu Leu Pro Asp Ile Asn Asp Gln Glu Phe Thr Arg Ile Cys Gln 165 170 175 Tyr Leu Leu Thr Glu Arg Phe Pro Phe Ser Ser Arg Gly Phe Phe Arg 185 Ile Met Val Arg Asp Cys Glu Ala Gly Met Val Asp Glu Asp Val Leu 195 200 205 200 Tyr Arg Phe Cys His Leu Pro Glu Phe Leu Tyr Val Arg Ser Leu Leu 215 220 Phe Gly Ala Glu Ile Glu Ala Ala Ser Val Ala Ser Leu Ala Arg Met 230 235 Ile Ile Gln Gly Glu Asp Leu Phe Phe Ser Leu Cys Cys Leu Glu 245 250 Asn Arg Gln Thr Ala Ile Ser Asp His Gln Arg Arg Cys Phe Leu Lys 260 265 270 265 270 Ala Tyr Val Asp Arg Gln Glu Pro Leu Ala Ala Leu Leu Leu Leu Val 275 280 285 275 280 His Asp Ala Asp Trp Val Leu His Glu Phe Ser Asp Ser Asp Leu Gln 295 Ser Phe Ile Gln Leu Leu Pro Arg Glu Ala His Tyr Thr Lys Lys Phe 310 315 Leu Gly Cys Val Ala Gln Ser Cys Arg Leu Gly Ile Leu Leu Glu Gly 325 330 <210> 573 <211> 426 <212> PRT <213> C. Trachomatis D serovar <400> 573 Met Tyr Val Arg Ser Ile Phe Phe Ser Ile Ile Ala Phe Leu Thr Val 10 Gly Cys Ser Phe Ser Pro Pro Glu Ser Gly Leu Ile Ile Ala Ile His Asp Asp Pro Arg Ser Leu Ser Pro Glu Lys Gly Glu Asn Ala Phe His 40 Phe Ser Leu Ser Lys Ala Leu Phe Ala Thr Leu Phe Arg Glu Glu Leu 55 Ser Gly Leu Thr Pro Ala Leu Val Ser Ser Tyr Gln Val Ser Glu Asp 75 Gly Arg Phe Tyr Arg Phe Cys Ile Arg Lys Asp Ala Lys Trp Ser Asp 8.5 90 Gly Ser Leu Leu Leu Ala Glu Asp Val Ile Ala Ala Trp Glu His Thr 100 105 Lys Gln Ala Gly Arg Tyr Ser Leu Leu Phe Glu Lys Leu Ser Phe Arg 120 125 Ala Ser Ser Ser Ser Glu Ile Leu Ile Glu Leu Lys Glu Pro Glu Pro 130 135 140 Gln Leu Leu Ala Ile Leu Ala Ser Pro Phe Phe Ala Val Tyr Arg Pro 150 155 Glu Asn Pro Phe Leu Ser Ser Gly Pro Phe Met Pro Lys Thr Tyr Val 170 Gln Gly Gln Thr Leu Val Leu Gln Lys Asn Pro Tyr Tyr Asp His 185 180 190

Ala His Val Glu Leu His Ser Ile Asp Phe Arg Ile Ile Pro Asn Ile

200 Tyr Thr Ala Leu His Leu Leu Arg Arg Gly Asp Val Asp Trp Val Gly 215 220 Gln Pro Trp His Gln Gly Ile Pro Phe Glu Leu Arg Thr Thr Ser Ala 230 235 Leu Tyr Thr His Tyr Ser Val Asp Gly Thr Phe Trp Leu Ile Leu Asn 245 250 255 Pro Lys Asp Pro Val Leu Ser Ser Leu Ser Asn Arg Gln Arg Leu Ile 260 . 265 Ala Ala Val Gln Lys Glu Lys Leu Val Lys Gln Ala Leu Gly Thr Gln 280 Tyr Arg Val Ala Glu Ser Ser Pro Ser Pro Glu Gly Ile Ile Ala His 290 295 300 Gln Glu Ala Ser Thr Pro Phe Pro Gly Lys Ile Thr Leu Ile Tyr Pro 310 315 Asn Asn Ile Thr Arg Cys Gln Arg Leu Ala Glu Val Leu Gln Glu Gln 325 330 Cys Arg Asp Ala Gly Ile Gln Leu Thr Leu Glu Gly Leu Glu Tyr His 345 350 Val Phe Val Gln Lys Arg Ala Thr Gln Asp Phe Ser Val Ser Thr Ala 355 360 365 Thr Ser Ile Ala Phe His Pro Leu Ala Lys Ser Lys Phe Asp Gln Thr 370 375 380Ala Leu Asp Asn Phe Thr Cys Leu Pro Leu Tyr His Ile Glu Tyr Asp 385 390 395 Tyr Ile Leu Ser Arg Pro Leu Asp Gln Ile Val His Tyr Pro Ser Gly 405 410 Ser Val Asp Leu Thr Tyr Ala His Phe His 420 <210> 574 <211> 605 <212> PRT <213> C. Trachomatis D serovar <400> 574 Met Gln Asn Ile Leu Arg Thr Ser Ser Cys Arg Tyr Met Phe Leu Leu 10 Gly Ile Arg Ser Val Trp Asn Arg Val Ala Val Val Asn Asn Phe Arg Gly Ser Ser Trp Lys Ile Val Ala Ile Pro Ser Cys Ile Leu Phe Thr 35 40 45

Leu Ile Phe His Leu Pro Arg Trp Leu Ile Asp Phe Gly Val Cys Thr 55 Asn Leu Ala Cys Ser Leu Ser Ile Ile Phe Trp Val Phe Ser Leu Arg 70 75 Ser Ser Ala Ser Ala Arg Ile Phe Pro Ser Leu Leu Tyr Leu Cys 85 90 Leu Leu Arg Leu Gly Leu Asn Leu Ala Ser Thr Arg Trp Ile Leu Ser 100 105 110 Ser Gly Trp Ala Ser Pro Leu Ile Phe Ala Leu Gly Asn Phe Phe Ser 115 120 Leu Gly Ser Ile Pro Val Ala Leu Thr Val Cys Leu Leu Leu Phe Leu 135 140 Val Asn Phe Leu Val Ile Thr Lys Gly Ala Glu Arg Ile Ala Glu Val 145 150 **155** Arg Ala Arg Phe Ser Leu Glu Ala Leu Pro Gly Lys Gln Met Ser Leu 165 170 Asp Ala Asp Ile Ala Ala Gly Arg Ile Gly Tyr Ser Arg Ala Ser Val 185 190 Lys Lys Ser Ser Leu Leu Glu Glu Ser Asp Tyr Phe Ser Ala Met Glu

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Leu Gly Val Asn Ile Leu Ala Ala Leu Phe Leu Gly Arg Ala Thr His
       230
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Val Gly Asp Leu Trp Leu Thr Val Leu Gly Asp Ala Leu Val Ser Gln 245 250 255
Ile Pro Ala Leu Leu Thr Ser Cys Ala Ala Ala Thr Leu Ile Ala Lys
        260 265 270
Val Gly Glu Lys Glu Ser Leu Ala Gln His Leu Leu Asp Tyr Tyr Glu
 275 280
                           285
Gln Ser Arg Gln Ser Phe Leu Phe Ile Ala Leu Ile Leu Cys Gly Met
                295
                                    300
Ala Cys Ile Pro Gly Ala Pro Lys Ala Leu Ile Leu Gly Phe Ser Val
               310
                               315
Leu Leu Phe Leu Gly Tyr Lys Asn Pro Ser Ser Gly Glu Thr Leu Leu
                            330
            325
                                             335
Phe Gln Lys Glu Arg Val Glu Phe Val Leu Pro Asp Glu Gly Val Gly
                 345
        340
                                          350
Asn Pro Ala Asn Leu Tyr Lys Asp Ala Arg Asn Gln Ile Tyr Gln Glu
   355 360
Leu Gly Val Val Phe Pro Glu Ala Ile Val Val Arg His Val Thr Gly 370 375 380
                                   380
Ser Ser Pro Arg Leu Ile Phe Ser Gly Gln Glu Val Ala Leu Arg Glu
              390
                              395
Leu Ser Cys Pro Ala Ile Leu Glu Ser Ile Arg Gln Leu Ala Pro Glu
         405
                           410
Thr Ile Ser Glu Arg Phe Val Thr Arg Leu Val Asp Glu Phe Arg Glu
         420
                         425
His Ala Phe Leu Ser Ile Glu Glu Ile Leu Pro Leu Lys Ile Ser Glu
           440
                                    445
Asn Ser Leu Ile Phe Leu Leu Arg Ala Leu Val Arg Glu Arg Val Ser
450 455 460
Leu His Leu Phe Pro Lys Ile Leu Glu Ala Ile Asp Val Tyr Gly Ser
             470
                                475
Gln Pro Lys Asn Ser Gln Glu Leu Val Glu Cys Val Arg Lys Tyr Leu
            485
                            490
Gly Lys Gln Ile Gly Leu Ser Leu Trp Asn Arg Gln Asp Val Leu Glu
         500
                         505
                                        510
Val Ile Thr Ile Asp Ser Leu Val Glu Gln Phe Val Arg Asp Ser Gln
 515
             520 525
Glu Lys Val Val Leu Asp Leu Asn Glu Lys Val Val Ala Gln Val Lys
                 535 540
His Leu Leu Arg Val Gly Glu Gly Asn Phe Arg Ala Ile Val Thr Gly 545 550 560
Ser Glu Thr Arg Lys Glu Leu Lys Arg Ile Val Asp Pro Tyr Phe Pro
            565
                           570
Asp Leu Leu Val Leu Ala His Ser Glu Leu Pro Glu Glu Ile Pro Ile
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Thr Leu Leu Gly Ala Val Ser Asp Glu Val Leu Leu Ser
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<400> 575

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<211> 173

<212> PRT

<213> C. Trachomatis D serovar

352

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Val Ile Ile Gly Gly Gln Ala Gly Ile Thr Gly His Ile Cys Ile Ala

353

275 280 285 Asp His Val Ile Met Met Ala Gln Thr Gly Val Thr Lys Ser Ile Thr 295 300 Ser Pro Gly Ile Tyr Gly Gly Ala Pro Ala Arg Pro Tyr Gln Glu Ile 310 315 His Arg Gln Val Ala Lys Val Arg Asn Leu Pro Arg Leu Glu Glu Arg 325 330 335 Ile Ala Ala Leu Glu Lys Leu Val Gln Lys Leu Glu Ala Leu Ser Glu 345 350 Gln His <210> 577 <211> 421 <212> PRT <213> C. Trachomatis D serovar <400> 577 Met Thr Ala Ser Gly Gly Ala Gly Gly Leu Gly Ser Thr Gln Thr Val 10 Asp Val Ala Arg Ala Gln Ala Ala Ala Ala Thr Gln Asp Ala Gln Glu Val Ile Gly Ser Gln Glu Ala Ser Glu Ala Ser Met Leu Lys Gly Cys Glu Asp Leu Ile Asn Pro Ala Ala Ala Thr Arg Ile Lys Lys Lys Gly 55 Glu Lys Phe Glu Ser Leu Glu Ala Arg Arg Lys Pro Thr Ala Asp Lys 70 75 Ala Glu Lys Lys Ser Glu Ser Thr Glu Glu Lys Gly Asp Thr Pro Leu Glu Asp Arg Phe Thr Glu Asp Leu Ser Glu Val Ser Gly Glu Asp Phe 105 Arg Gly Leu Lys Asn Ser Phe Asp Asp Ser Ser Pro Asp Glu Ile 120 125 Leu Asp Ala Leu Thr Ser Lys Phe Ser Asp Pro Thr Ile Lys Asp Leu 135 140 Ala Leu Asp Tyr Leu Ile Gln Thr Ala Pro Ser Asp Gly Lys Leu Lys 150 155 Ser Thr Leu Ile Gln Ala Lys His Gln Leu Met Ser Gln Asn Pro Gln 165 170 175 Ala Ile Val Gly Gly Arg Asn Val Leu Leu Ala Ser Glu Thr Phe Ala 180 185 190 Ser Arg Ala Asn Thr Ser Pro Ser Ser Leu Arg Ser Leu Tyr Phe Gln 200 205 Val Thr Ser Ser Pro Ser Asn Cys Ala Asn Leu His Gln Met Leu Ala 215 Ser Tyr Leu Pro Ser Glu Lys Thr Ala Val Met Glu Phe Leu Val Asn 230 235 Gly Met Val Ala Asp Leu Lys Ser Glu Gly Pro Ser Ile Pro Pro Ala 245 250 Lys Leu Gln Val Tyr Met Thr Glu Leu Ser Asn Leu Gln Ala Leu His 260 265 Ser Val Asn Ser Phe Phe Asp Arg Asn Ile Gly Asn Leu Glu Asn Ser 280 Leu Lys His Glu Gly His Ala Pro Ile Pro Ser Leu Thr Thr Gly Asn 295 300 Leu Thr Lys Thr Phe Leu Gln Leu Val Glu Asp Lys Phe Pro Ser Ser 310 315 Ser Lys Ala Gln Lys Ala Leu Asn Glu Leu Val Gly Pro Asp Thr Gly

330

Pro Gln Thr Glu Val Leu Asn Leu Phe Phe Arg Ala Leu Asn Gly Cys

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340
                              345
Ser Pro Arg Ile Phe Ser Gly Ala Glu Lys Lys Gln Gln Leu Ala Ser 355 360 365
Val Ile Thr Asn Thr Leu Asp Ala Ile Asn Ala Asp Asn Glu Asp Tyr
                    375
                                380
Pro Lys Pro Gly Asp Phe Pro Arg Ser Ser Phe Ser Ser Thr Pro Pro
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His Ala Pro Val Pro Gln Ser Glu Ile Pro Thr Ser Pro Thr Ser Thr
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                                 410
Gln Pro Pro Ser Pro
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<213> C. Trachomatis D serovar
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Gln Asn Lys His Met Leu Glu His Thr Phe Tyr Val Lys Trp Ser Lys
     20
                               25
Gly Glu Leu Thr Lys Glu Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr
                          40
Leu His Ile Lys Ala Phe Pro Lys Tyr Leu Ser Ala Ile His Ser Arg 50 55
Cys Asp Asp Leu Glu Ala Arg Lys Leu Leu Leu Asp Asn Leu Met Asp
              70
                                   75
Glu Glu Asn Gly Tyr Pro Asn His Ile Asp Leu Trp Lys Gln Phe Val
              85
                                  90
Phe Ala Leu Gly Val Thr Pro Glu Glu Leu Glu Ala His Glu Pro Ser
           100
                              105
                                               110
Glu Ala Ala Lys Ala Lys Val Ala Thr Phe Met Arg Trp Cys Thr Gly
115 120 125
                          120
                                              125
Asp Ser Leu Ala Ala Gly Val Ala Ala Leu Tyr Ser Tyr Glu Ser Gln
130 135 140
Ile Pro Arg Ile Ala Arg Glu Lys Ile Arg Gly Leu Thr Glu Tyr Phe 145 150 155 160
Gly Phe Ser Asn Pro Glu Asp Tyr Ala Tyr Phe Thr Glu His Glu Glu 165 170 175
Ala Asp Val Arg His Ala Arg Glu Glu Lys Ala Leu Ile Glu Met Leu
         180
                              185
                                     190
Leu Lys Asp Asp Ala Asp Lys Val Leu Glu Ala Ser Gln Glu Val Thr
195 200 205
Gln Ser Leu Tyr Gly Phe Leu Asp Ser Phe Leu Asp Pro Gly Thr Cys
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Cys Ser Cys His Gln Ser Tyr
<210> 579
<211> 243
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<400> 579
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Leu Gln Glu Ile Leu Gln Glu Ala Leu Pro Pro Leu Gln Glu Arg Ser
                               25
                                                 30
Val Val Val Ser Ser Lys Ile Val Ser Leu Cys Glu Gly Ala Val
                   40
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Ala Asp Ala Arg Met Cys Lys Ala Glu Leu Ile Lys Lys Glu Ala Asp 55 Ala Tyr Leu Phe Cys Glu Lys Ser Gly Ile Tyr Leu Thr Lys Lys Glu 75 Gly Ile Leu Ile Pro Ser Ala Gly Ile Asp Glu Ser Asn Thr Asp Gln 85 90 Pro Phe Val Leu Tyr Pro Lys Asp Ile Leu Gly Ser Cys Asn Arg Ile 100 \$105\$Gly Glu Trp Leu Arg Asn Tyr Phe Arg Val Lys Glu Leu Gly Val Ile 120 Ile Thr Asp Ser His Thr Thr Pro Met Arg Arg Gly Val Leu Gly Ile 135 140 Gly Leu Cys Trp Tyr Gly Phe Ser Pro Leu His Asn Tyr Ile Gly Ser 150 155 Leu Asp Cys Phe Gly Arg Pro Leu Gln Met Thr Gln Ser Asn Leu Val 165 170 Asp Ala Leu Ala Val Ala Ala Val Val Cys Met Gly Glu Gly Asn Glu 180 185 190 Gln Thr Pro Leu Ala Val Ile Glu Gln Ala Pro Asn Met Val Tyr His 195 . 200 205 Ser His Pro Thr Ser Arg Glu Glu Tyr Cys Ser Leu Arg Ile Asp Glu 215 220 Thr Glu Asp Leu Tyr Gly Pro Phe Leu Gln Ala Val Thr Trp Ser Gln 230 Glu Lys Lys <210> 580 <211> 383 <212> PRT <213> C. Trachomatis D serovar <400> 580 Met Leu Pro His Gln Gln Asn Ser Ser Ser Glu Arg Ala Arg His His 1 5 10 Glu Ser Arg Ser His Arg His Ser Ser Ser Ser Arg His His Val Thr 25 Arg Ser Gln Ser Ser Ala Leu Pro Gln Leu Gln Glu Arg Pro Val Pro 35 40 4.5 His Pro Leu Ala Glu Arg Glu Leu Ile Ile Phe His Ser Val His Gln 50 55 60 Gln Gln Asn Asn Pro Leu Arg Met Ile Cys Asp Thr Ile Arg Gln 70 75 Ala Gln Arg Gly Ile Phe Met Arg Ile Tyr Thr Ile Ser Ser Asp Asp 90 Ile Ile Gln Ser Leu Ile Gln Thr Ser His His Val Pro Val Glu Val 100 105 110 Lys Tyr His Cys Gly Glu Ser Leu Pro Val Ala Cys Gln Asn Ser Arg 115 120

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Ser Ile Phe Val Leu Met Tyr Ile Phe Leu Ser Pro Glu Phe Phe Leu
                  230
Ala Leu Ala Gln Ala Met Arg Arg Gly Val Arg Val Lys Val Ile Ile
               245
                                  250
Asp Asn His Ser Lys Gln Asp Thr Cys Lys Leu Leu Ser Lys Leu Gly
           260
                              265
                                                270
Ile Gln Leu Pro Ile Tyr Glu Arg Lys Thr Glu Gly Val Leu His Thr
                          280
Lys Ile Cys Cys Ile Asp Asn Lys Thr Leu Ile Phe Gly Ser Ala Asn
                       295
                                          300
Trp Ser Gly Ala Gly Met Ile Lys Asn Phe Glu Asp Leu Phe Ile Leu
                  310
                             315
Arg Pro Ile Thr Glu Thr Gln Leu Gln Ala Phe Met Asp Val Trp Ser 325 330 335
Leu Leu Glu Thr Asn Ser Ser Tyr Leu Ser Pro Glu Ser Val Leu Thr
                              345
Ala Pro Thr Pro Ser Ser Arg Pro Thr Gln Gln Asp Thr Asp Ser Asp
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                                              365
Asp Glu Gln Pro Ser Thr Ser Gln Gln Asp Ile Arg Met Arg Lys
                      375
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<213> C. Trachomatis D serovar
Met Trp Phe Phe Leu Gly Ser Pro Ser Ala Ile Thr Asn Phe Ser Arg
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Val Asp Val Ala Leu Asn Leu Arg Ile Asn Arg Gln Ile Arg Ala Pro
        20
                               25
Arg Val Arg Val Ile Gly Ser Ala Gly Glu Gln Leu Gly Ile Leu Ser
      35
                           40
Ile Lys Glu Ala Leu Asp Leu Ala Lys Glu Ala Asn Leu Asp Leu Val
Glu Val Ala Ser Asn Ser Glu Pro Pro Val Cys Lys Ile Met Asp Tyr
                 70
                                      75
Gly Lys Tyr Arg Tyr Asp Val Thr Lys Lys Glu Lys Asp Ser Lys Lys 85 90 95
               85
                                  90
                                                      95
Ala Gln His Gln Val Arg Ile Lys Glu Val Lys Leu Lys Pro Asn Ile
          100
                              105
Asp Asp Asn Asp Phe Leu Thr Lys Ala Lys Gln Ala Arg Ala Phe Ile
      115
                         120
                                              125
Glu Lys Gly Asn Lys Val Lys Val Ser Cys Met Phe Arg Gly Arg Glu
130 135
Leu Ala Tyr Pro Glu His Gly Tyr Lys Val Ile Gln Arg Met Cys Gln
145 150 155 160
                  150
                                   155
Gly Leu Glu Asp Ile Gly Phe Val Glu Ser Glu Pro Lys Leu Asn Gly
              165
                                 170
Arg Ser Leu Ile Cys Val Ile Ala Pro Gly Thr Leu Lys Thr Lys Lys
Lys
<210> 582
<211> 264
<212> PRT
<213> C. Trachomatis D serovar
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Met Gly Asn Ser Gly Phe Tyr Leu Tyr Asn Thr Glu Asn Cys Val Phe

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195 200 205 Thr Ser Ile Thr Asn Thr Gly Leu Thr Pro Thr Thr Tyr Ser Leu Arg 210 215 220 Val Gly Gly Leu Glu Ser Gly Val Val Trp Val Asn Ala Leu Ser Asn 225 230 235 Gly Asn Asp Ile Leu Gly Ile Thr Asn Thr Ser Asn Val Ser Phe Leu 245 250 Glu Val Ile Pro Gln Thr Asn Ala 260

<210> 583

<211> 1053

<212> PRT

<213> C. Trachomatis D serovar

Met Phe Thr Arg Ile Val Met Val Asp Leu Gln Glu Lys Gln Cys Thr 10 Ile Val Lys Arg Asn Gly Met Phe Val Pro Phe Asp Arg Asn Arg Ile 20 Phe Gln Ala Leu Glu Ala Ala Phe Arg Asp Thr Arg Arg Ile Asp Asp 40 His Met Pro Leu Pro Glu Asp Leu Glu Ser Ser Ile Arg Ser Ile Thr 55 His Gln Val Val Lys Glu Val Val Gln Lys Ile Thr Asp Gly Gln Val 65 70 75 80Val Thr Val Glu Arg Ile Gln Asp Met Val Glu Ser Gln Leu Tyr Val 85 90 Asn Gly Leu Gln Asp Val Ala Arg Asp Tyr Ile Val Tyr Arg Asp Asp 100 105 110 Arg Lys Ala His Arg Lys Lys Ser Trp Gln Ser Leu Ser Val Val Arg 115 120 125 Arg Cys Gly Thr Val Val His Phe Asn Pro Met Lys Ile Ser Ala Ala 130 135 140 Leu Glu Lys Ala Phe Arg Ala Thr Asp Lys Thr Glu Gly Met Thr Pro 150 155 Ser Ser Val Arg Glu Glu Ile Asn Ala Leu Thr Gln Asn Ile Val Ala

														175	
				165	_	D	01	63	170			~7 -	7	175	C1
Glu	Ile	Glu	Glu 180	Cys	Cys	Pro	GIn	185	Asp	Arg	Arg	TTE	190	Ile	GIU
Lys	Ile	Gln 195	Asp	Ile	Val	Glu	Gln 200	Gln	Leu	Met	Val	Val 205	Gly	His	Tyr
Ala	Val 210	Ala	Lys	Asn	Tyr	Ile 215	Leu	Tyr	Arg	Glu	Ala 220	Arg	Ala	Arg	Val
Arg 225		Asn	Arg	Glu	Glu 230	Asp	Gly	Ser	Thr	Glu 235	Lys	Thr	Ile	Ala	Glu 240
	Ala	Val	Glu	Val 245	Leu	Ser	Lys	Asp	Gly 250	Ser	Thr	Tyr	Thr	Met 255	Thr
His	Ser	Gln	Leu 260	Leu	Ala	His	Leu	Ala 265	Arg	Ala	Суѕ	Ser	Arg 270	Phe	Pro
Glu	Thr	Thr 275	Asp	Ala	Ala	Leu	Leu 280	Thr	Asp	Met	Ala	Phe 285	Ala	Asn	Phe
Tyr	Ser 290	Gly	Ile	Lys	Glu	Ser 295	Glu	Val	Val	Leu	Ala 300	Cys	Ile	Met	Ala
Ala 305	Arg	Ala	Asn	Ile	Glu 310	Lys	Glu	Pro	Asp	Tyr 315	Ala	Phe	Val	Ala	Ala 320
				325					330					Ser 335	
_			340					345					350	Arg	
		355					360					365		His	
	370		_			375					380			Asp	
385			_		390					395				Phe	400
			-	405			•		410					Met 415	
			420					425					430	Trp	
		435	-				440			-		445		Ala	
	450					455					460			Ser	
465			,		470					475				Val	480
	_			485					490					Asn 495	
_			500					505					510	Asn	
		515					520					525		Thr	
	530					535					540			Val	_
545					550					555				Arg	560
		_	_	565					570					Ala 575	
-			580					585					590	Thr	
		595					600					605		Tyr	
	610					615					620			Asp	
625					630					635				Trp	640
тЛг	меt	ьeu	ser	мет 645		5116	оти	TUL	650	птз	LTO	тър	net	Thr 655	THE

Lys Asp Pro Ser Asn Ile Arg Ser Ala Gln Asp His Lys Gly Val Val

665

660

359

Arg Cys Ser Asn Leu Cys Thr Glu Ile Leu Leu Asn Cys Ser Glu Thr 680 685 Glu Thr Ala Val Cys Asn Leu Gly Ser Ile Asn Leu Val Gln His Ile 695 Val Gly Asp Gly Leu Asp Glu Glu Lys Leu Ser Glu Thr Ile Ser Ile 705 710 715 720 Ala Val Arg Met Leu Asp Asn Val Ile Asp Ile Asn Phe Tyr Pro Thr 725 730 735 Lys Glu Ala Lys Glu Ala Asn Phe Ala His Arg Ala Ile Gly Leu Gly 740 745 Val Met Gly Phe Gln Asp Ala Leu Tyr Lys Leu Asp Ile Ser Tyr Ala 755 760 Ser Gln Glu Ala Val Glu Phe Ala Asp Tyr Ser Ser Glu Leu Ile Ser 775 780 Tyr Tyr Ala Ile Gln Ala Ser Cys Leu Leu Ala Lys Glu Arg Gly Thr 785 790 795 800 Tyr Ser Ser Tyr Lys Gly Ser Lys Trp Asp Arg Gly Leu Leu Pro Ile 805 810 815 Asp Thr Ile Gln Leu Leu Ala Asn Tyr Arg Gly Glu Ala Asn Leu Gln 825 Met Asp Thr Ser Ser Arg Lys Asp Trp Glu Pro Ile Arg Ser Leu Val 835 840 845 Lys Glu His Gly Met Arg His Cys Gln Leu Met Ala Ile Ala Pro Thr 855 860 Ala Thr Ile Ser Asn Ile Ile Gly Val Thr Gln Ser Ile Glu Pro Thr 870 875 880 Tyr Lys His Leu Phe Val Lys Ser Asn Leu Ser Gly Glu Phe Thr Ile 885 890 895 Pro Asn Val Tyr Leu Ile Glu Lys Leu Lys Leu Gly Ile Trp Asp 900 905 910 Ala Asp Met Leu Asp Asp Leu Lys Tyr Phe Asp Gly Ser Leu Leu Glu 920 925 Ile Glu Arg Ile Pro Asp His Leu Lys His Ile Phe Leu Thr Ala Phe 935 940 Glu Ile Glu Pro Glu Trp Ile Ile Glu Cys Ala Ser Arg Arg Gln Lys 945 950 955 Trp Ile Asp Met Gly Gln Ser Leu Asn Leu Tyr Leu Ala Gln Pro Asp 965 970 975 Gly Lys Lys Leu Ser Asn Met Tyr Leu Thr Ala Trp Lys Lys Gly Leu 980 985 990 980 985 Lys Thr Thr Tyr Tyr Leu Arg Ser Ser Ser Ala Thr Thr Val Glu Lys 1000 1005 Ser Phe Val Asp Ile Asn Lys Arg Gly Ile Gln Pro Arg Trp Met Lys 1015 1020 Asn Lys Ser Ala Ser Ala Gly Ile Ile Val Glu Arg Ala Lys Lys Ala 1025 1030 1035 Pro Val Cys Ser Leu Glu Glu Gly Cys Glu Ala Cys Gln 1045 <210> 584 <211> 346 <212> PRT <213> C. Trachomatis D serovar Met Gln Ala Asp Ile Leu Asp Gly Lys Gln Lys Arg Val Asn Leu Asn 10

Ser Lys Arg Leu Val Asn Cys Asn Gln Val Asp Val Asn Gln Leu Val

360

Pro Ile Lys Tyr Lys Trp Ala Trp Glu His Tyr Leu Asn Gly Cys Ala 35 40 45Asn Asn Trp Leu Pro Thr Glu Ile Pro Met Gly Lys Asp Ile Glu Leu Trp Lys Ser Asp Arg Leu Ser Glu Asp Glu Arg Arg Val Ile Leu Leu 75 Asn Leu Gly Phe Phe Ser Thr Ala Glu Ser Leu Val Gly Asn Asn Ile 85 90 Val Leu Ala Ile Phe Lys His Val Thr Asn Pro Glu Ala Arg Gln Tyr 100 ' 105 Leu Leu Arg Gln Ala Phe Glu Glu Ala Val His Thr His Thr Phe Leu 120 125 Tyr Ile Cys Glu Ser Leu Gly Leu Asp Glu Lys Glu Ile Phe Asn Ala 135 140 Tyr Asn Glu Arg Ala Ala Ile Lys Ala Lys Asp Asp Phe Gln Met Glu 145 150 155 160 Ile Thr Gly Lys Val Leu Asp Pro Asn Phe Arg Thr Asp Ser Val Glu 165 · 170 175 Gly Leu Gln Glu Phe Val Lys Asn Leu Val Gly Tyr Tyr Ile Ile Met 185 Glu Gly Ile Phe Phe Tyr Ser Gly Phe Val Met Ile Leu Ser Phe His 200 195 205 Arg Gln Asn Lys Met Ile Gly Ile Gly Glu Gln Tyr Gln Tyr Ile Leu 210 215 220 Arg Asp Glu Thr Ile His Leu Asn Phe Gly Ile Asp Leu Ile Asn Gly 230 235 Ile Lys Glu Glu Asn Pro Glu Ile Trp Thr Pro Glu Leu Gln Glu 250 Ile Val Glu Leu Ile Lys Arg Ala Val Asp Leu Glu Ile Glu Tyr Ala 265 270 Gln Asp Cys Leu Pro Arg Gly Ile Leu Gly Leu Arg Ala Ser Met Phe 275 280 285 Ile Asp Tyr Val Gln His Ile Ala Asp Arg Arg Leu Glu Arg Ile Gly 295 300 Leu Lys Pro Ile Tyr His Thr Lys Asn Pro Phe Pro Trp Met Ser Glu 310 315 320 Thr Ile Asp Leu Asn Lys Glu Lys Asn Phe Phe Glu Thr Arg Val Ile 325 330 335 Glu Tyr Gln His Ala Ala Ser Leu Thr Trp 340 <210> 585 <211> 326 <212> PRT <213> C. Trachomatis D serovar <400> 585 Met Ser Phe Phe His Thr Arg Lys Tyr Lys Leu Ile Leu Arg Gly Leu 10 Leu Cys Leu Ala Gly Cys Phe Leu Met Asn Ser Cys Ser Ser Ser Arg 25 Gly Asn Gln Pro Ala Asp Glu Ser Ile Tyr Val Leu Ser Met Asn Arg Met Ile Cys Asp Cys Val Ser Arg Ile Thr Gly Asp Arg Val Lys Asn 55 60

Asn Pro Lys Val Val Asp Leu Gly Gln Arg Leu Leu Asn Lys Asn Cys 115 120 Phe Asp Leu Leu Ser Glu Glu Gly Phe Pro Asp Pro His Ile Trp Thr 135 130 140 Asp Met Arg Val Trp Gly Ala Ala Val Lys Glu Met Ala Ala Ala Leu 150 155 Ile Gln Gln Phe Pro Gln Tyr Glu Glu Asp Phe Gln Lys Asn Ala Asp 165 170 Gln Ile Leu Ser Glu Met Glu Glu Leu Asp Arg Trp Ala Ala Arg Ser 180 185 190 Leu Ser Thr Ile Pro Glu Lys Asn Arg Tyr Leu Val Thr Gly His Asn 200 . 205 Ala Phe Ser Tyr Phe Thr Arg Arg Tyr Leu Ser Ser Asp Ala Glu Arg 215 220 Val Ser Gly Glu Trp Arg Ser Arg Cys Ile Ser Pro Glu Gly Leu Ser 230 235 Pro Glu Ala Gln Ile Ser Ile Arg Asp Ile Met Arg Val Val Glu Tyr 250 Ile Ser Ala Asn Asp Val Glu Val Val Phe Leu Glu Asp Thr Leu Asn 265 270 Gln Asp Ala Leu Arg Lys Ile Val Ser Cys Ser Lys Ser Gly Gln Lys 280 Ile Arg Leu Ala Lys Ser Pro Leu Tyr Ser Asp Asn Val Cys Asp Asn 295 Tyr Phe Ser Thr Phe Gln His Asn Val Arg Thr Ile Thr Glu Glu Leu 310 Gly Gly Thr Val Leu Glu 325 <210> 586 <211> 102 <212> PRT <213> C. Trachomatis D serovar <400> 586 Met Gln Asn Lys Arg Lys Val Arg Asp Asp Phe Ile Lys Ile Val Lys Asp Val Lys Lys Asp Phe Pro Glu Leu Asp Leu Lys Ile Arg Val Asn Lys Glu Lys Val Thr Phe Leu Asn Ser Pro Leu Glu Leu Tyr His Lys 35 40 Ser Val Ser Leu Ile Leu Gly Leu Leu Gln Gln Ile Glu Asn Ser Leu 55 Gly Leu Phe Pro Asp Ser Pro Val Leu Glu Lys Leu Glu Asp Asn Ser 70 75 Leu Lys Leu Lys Lys Ala Leu Ile Met Leu Ile Leu Ser Arg Lys Asp 85 90 Met Phe Ser Lys Ala Glu 100 <210> 587 <211> 243 <212> PRT <213> C. Trachomatis D serovar <400> 587 Val Gly Cys Asn Leu Ala Gln Phe Leu Gly Lys Lys Val Leu Leu Ala 10 Asp Leu Asp Pro Gln Ser Asn Leu Ser Ser Gly Leu Gly Ala Ser Val 20 Arg Asn Asn Gln Lys Gly Leu His Asp Ile Val Tyr Lys Ser Asn Asp

362

Leu Lys Ser Ile Ile Cys Glu Thr Lys Lys Asp Ser Val Asp Leu Ile 55 60 Pro Ala Ser Phe Leu Ser Glu Gln Phe Arg Glu Leu Asp Ile His Arg 70 Gly Pro Ser Asn Asn Leu Lys Leu Phe Leu Asn Glu Tyr Cys Ala Pro 90 Phe Tyr Asp Ile Cys Ile Ile Asp Thr Pro Pro Ser Leu Gly Gly Leu 100 105 110 Thr Lys Glu Ala Phe Val Ala Gly Asp Lys Leu Ile Ala Cys Leu Thr 115 120 Pro Glu Pro Phe Ser Ile Leu Gly Leu Gln Lys Ile Arg Glu Phe Leu 135 140 Ser Ser Val Gly Lys Pro Glu Glu Glu His Ile Leu Gly Ile Ala Leu 150 155 Ser Phe Trp Asp Asp Arg Asn Ser Thr Asn Gln Met Tyr Ile Asp Ile
165 170 175 165 170 Ile Glu Ser Ile Tyr Lys Asn Lys Leu Phe Ser Thr Lys Ile Arg Arg 185 Asp Ile Ser Leu Ser Arg Ser Leu Leu Lys Glu Asp Ser Val Ala Asn 195 200 205 Val Tyr Pro Asn Ser Arg Ala Ala Glu Asp Ile Leu Lys Leu Thr His 210 215 220 Glu Ile Ala Asn Ile Leu His Ile Glu Tyr Glu Arg Asp Tyr Ser Gln 230 235 Arg Thr Thr

<210> 588

<211> 527

· <212> PRT

<213> C. Trachomatis D serovar

Met Pro Ser Leu Ser Gln Ser Arg Arg Ile Ile Gln Gln Ser Ser Ile 10 Arg Lys Ile Trp Asn Gln Ile Asp Thr Ser Pro Lys His Gly Val Cys 20 Val Pro Leu Phe Ser Leu Tyr Thr Gln Glu Ser Cys Gly Ile Gly Glu 40 Phe Leu Asp Leu Ile Pro Met Ile Asp Trp Cys Ile Ser Cys Gly Phe 50 55 60 Gln Ile Leu Gln Ile Leu Pro Ile Asn Asp Thr Gly Ser Cys Ser Ser 70 75 Pro Tyr Asn Ser Ile Ser Ser Ile Ala Leu Asn Pro Leu His Leu Ser 85 90 Ile Ser Ala Leu Pro Tyr Lys Glu Glu Val Pro Ala Ala Glu Thr Arg 100 105 110 Ile Arg Glu Met Gln Gln Leu Ser Gln Leu Pro Gln Val His Tyr Glu 115 120 125 Lys Val Arg Ser Met Lys Arg Asp Phe Phe Gln Glu Tyr Tyr Arg Val 135 140 Cys Lys Gln Lys Lys Leu Thr Asp His Pro Asp Phe Tyr Ala Phe Cys 155 160 Glu Gln Glu Lys Tyr Trp Leu His Pro Tyr Ala Leu Phe Arg Ser Ile 170 Arg Glu His Leu Asp Asn Leu Pro Ile Asn His Trp Pro Thr Thr Tyr 180 185 Thr Asp Leu Ser Gln Ile Thr Glu His Glu Arg Thr Phe Ala Glu Asp 200 Ile Gln Phe His Ser Tyr Leu Gln Tyr Leu Cys Phe Gln Gln Met Thr

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215
Gln Val Arg Glu His Ala Asn Cys Lys Ser Cys Leu Ile Lys Gly Asp
         230
                          235
Ile Pro Ile Leu Ile Ser Lys Asp Ser Cys Asp Val Trp Phe Tyr Arg
            245
                              250
His Tyr Phe Ser Ser Ser Glu Ser Val Gly Ala Pro Pro Asp Leu Tyr
         260
                           265
                                             270
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Pro Ser Asp Val Lys Arg Met Leu Glu Ser Phe Ala Val Cys Gly Thr
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Pro Phe Asp Gln Tyr Asp Pro Leu Ser Val Thr Ser Leu Ser Thr His
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Asp Ser Ser Thr Leu Ala Ser Trp Trp Lys Glu Ser Pro Gln Glu Ser
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Lys Leu Phe Ala Gln Phe Leu Gly Leu Pro Tyr Ser Ser Thr Leu Ser
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Ile Phe Arg Ile Asn Leu Ile Asn Asp Tyr Leu Ala Leu Phe Pro Asp
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Leu Ile Ser Lys Thr Pro Arg Tyr Glu Arg Ile Asn Leu Pro Gly Thr
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Gly Tyr Asn Asn Val Ala Val Gln Ile Glu Glu Asp Gly Asn Ser Gly
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Arg Gln Lys Ile Phe Lys Ala Ala Leu Ser Ile Asn Gly Ser Pro Gln
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Ser Asn Ile Lys Gly Thr Leu Gly Tyr Gly Glu Ile Ser Asn Gln Leu
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Tyr Leu Cys Asp Arg Leu Asn Met Thr Tyr Leu Asn Gly Glu Lys Leu
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WO 02/08267 364

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395

390

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Leu Val Leu Tyr Thr Asp Gly Val Thr Glu Ala Ser Asn Lys His Gly 580 585 590
Glu Met Phe Gly Glu Glu Arg Leu Lys Ala Leu Val Ala Ser Leu Thr
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Asp Pro Phe Asn Asp Glu Asn Gly Asn Ile Leu Ser Ser Glu Thr Leu
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Ala Leu Leu Lys Asn Arg Tyr Gly Leu Asp Lys Pro Leu Phe Thr Gln
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Tyr Leu Ile Tyr Leu Lys Cys Leu Leu Thr Leu Asp Phe Gly Glu Ser 65 70 75 80
Leu Ile Tyr Lys Asp Arg Thr Val Ile Ser Ile Ile Ala Ala Ala Leu
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Pro Ser Ser Ala Ile Leu Gly Leu Glu Ser Leu Cys Leu Ser Leu Phe
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Gly Gly Ile Thr Leu Gly Ile Leu Ala Ala Phe Tyr Lys Lys Ser Cys 115 120 125
Gly Arg Thr Ile Phe Phe Ser Ser Val Ile Gln Ile Ser Val Pro Ala
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Phe Val Ile Gly Ala Phe Leu Gln Tyr Val Phe Ala Ile Lys Tyr Ser
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Cys Leu Pro Ile Ala Cys Trp Gly Asn Phe Ser His Thr Leu Leu Pro
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366

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Thr Ile Thr Asn Thr Cys Ile Val Ala Glu Arg Cys His Leu Glu Leu Asp Phe Glu Thr Lys His Tyr Pro Ile Tyr Val Pro Glu Ala Leu Gln Lys Lys Gly Ser Tyr Thr Glu Glu Glu Arg Tyr Lys Ala Ser Ser Ala Phe Leu Glu Glu Leu Cys Glu Gln Gly Leu Thr Ser Lys Tyr Thr Pro 340 345 350 Glu Leu Cly His Ile Ala Lys Lys Phe Pro Gly Glu Asp Pro Leu 355 360 365 Thr Leu Val Lys Glu Arg Leu Lys Leu Glu Ser Ser Ile Ile Ile Ser Lys Gly Met Cys Asp Tyr Leu Leu Ile Val Trp Asp Ile Ile Asn Trp Ala Lys Asp His Gly Ile Pro Val Gly Pro Gly Arg Gly Ser Gly Ala Gly Ser Val Met Leu Phe Leu Leu Gly Ile Thr Glu Ile Glu Pro Ile Arg Phe Asp Leu Phe Phe Glu Arg Phe Ile Asn Pro Glu Arg Ile Ser Tyr Pro Asp Ile Asp Ile Asp Ile Cys Met Ile Gly Arg Glu Arg Val Ile Asn Tyr Ala Ile Glu Arg His Gly Lys Asp Asn Val Ala Gln Ile Ile Thr Phe Gly Thr Met Lys Ala Lys Met Ala Ile Lys Asp Val Gly Arg Thr Leu Asp Thr Pro Leu Ala Lys Val Asn Phe Ile Ala Lys His 500 505 510 Ile Pro Asp Leu Asn Ala Thr Ile Thr Ser Ala Leu Glu Ala Asp Pro Glu Leu Arg Gln Leu Tyr Val Asp Asp Ala Glu Ala Ala Glu Val Ile Asp Met Ala Lys Lys Leu Glu Gly Ser Ile Arg Asn Thr Gly Val His Ala Ala Gly Val Ile Ile Cys Gly Asp Pro Leu Thr Asn His Ile Pro Ile Cys Val Pro Lys Asp Ser Ser Met Ile Ser Thr Gln Tyr Ser Met Lys Pro Val Glu Ser Val Gly Met Leu Lys Val Asp Phe Leu Gly Leu
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Ala	Leu	Tyr 355	Val	Glu	Gly	Asp	Ile 360		Phe	Gln	Asp	Leu 365		Glu	Ile
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Ser	Ser	Pro	Gln	Ser 405	Gly	Ser	Gly	Ala	Thr 410	Thr	Val	Ser	Asn	Ser 415	
Asp	Ser	Ser	Ser 420	Gly	Ser	Asp	Ser	Asp 425	Thr	Ser	Glu	Thr	Val 430	Pro	Ala
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OTÀ	770		3	~ - y	<u> </u>	775		AT a	270	T11T	780	NOII	TTE	GET	LOII

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•	Ile 1010)				1015	5				1020)			
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Gln	Asn							,							
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Ser Gly Leu Val Thr 118	1090 Thr 5 Ser Asp Ser His 1170 Leu	Thr Gly Thr Leu 1159 Thr O Asp	Gln Asn Pro 1140 Gln 5 Ser Ile Ser	Ser Ile 1125 Ala Ala Thr Asn Ser 1205 His	Ser 1110 Thr Ser Ala Lys Lys 1190 Glu	1095 Gln Phe Lys Lys 1175 Glu Leu	Thr Ser Phe Gly 1160 Ile Glu	Gly Asp Asn Cys 1145 Lys Gly Asn Glu	Ala Asn 1130 Ser Thr Ser Ser Asn 1210 Val	Ile 1115 Ser) Ile Ile Thr Asn 1195 Lys	1100 Leu Leu Ala Ser Gln 1180 Pro	Thr Gln Gly Phe 1165 Asn Tyr	Gly Leu Asn Tyr 1150 Phe Val Thr	Asn 1135 Val) Asp Tyr Gly Pro 1215 Thr	Ala 1120 Gln Lys Cys Glu Thr 1200 Gln
Ser Gly Leu Val Thr 118 Ile	1090 Thr 5 Ser Asp Ser His 1170 Leu 5	Thr Gly Thr Leu 1159 Thr Asp Phe	Gln Pro 1140 Gln Ser Ile Ser Leu 1220 Val	Ser Ile 1125 Ala Ala Thr Asn Ser 1205 His	Ser 1110 Thr Ser Ala Lys Lys 1190 Glu Asn	1095 Gln Phe Lys Lys 1175 Glu Leu Gly	Thr Ser Phe Gly 1160 Ile Glu His	Gly Asp Asn Cys 1145 Lys Gly Asn Glu Leu 1225 Lys	Ala Asn 1130 Ser Thr Ser Ser Asn 1210 Val	Ile 1115 Ser) Ile Ile Thr Asn 1195 Lys)	1100 Leu Leu Ala Ser Gln 1180 Pro Ser Lys	Thr Gln Gly Phe 1165 Asn Tyr Tyr Glu	Leu Asn Tyr 1150 Phe Val Thr Lys 1230 Leu	Asn 1135 Val) Asp Tyr Gly Pro 1215 Thr	Ala 1120 Gln Lys Cys Glu Thr 1200 Gln Gln
Ser Gly Leu Val Thr 118 Ile Asn Leu	1090 Thr 5 Ser Asp Ser His 1170 Leu 5 Val	Thr Gly Thr Leu 1159 Thr Asp Phe Ile Val 1239 Gly	Gln Asn Pro 1140 Gln Ser Ile Ser Leu 1220 Val	Ser Ile 1125 Ala Ala Thr Asn Ser 1205 His	Ser 1110 Thr Ser Ala Lys 1190 Glu Asn	1095 Gln Phe Lys Lys 1175 Glu Leu Gly Glu	Thr Ser Phe Gly 1160 Ile Glu His Thr Gln 1240 Asn	Gly Asp Asn Cys 1145 Lys Gly Asn Glu Leu 1225 Lys	Ala Asn 1130 Ser Thr Ser Asn 1210 Val	Ile 1115 Ser Ile Ile Thr Asn 1195 Lys Leu	1100 Leu Leu Ala Ser Gln 1180 Pro Ser Lys	Thr Gln Gly Phe 1165 Asn Tyr Tyr Glu Lys 1245 Asn	Leu Asn Tyr 1150 Phe Val Thr Lys 1230 Leu	Asn 1135 Val) Asp Tyr Gly Pro 1215 Thr	Ala 1120 Gln Lys Cys Glu Thr 1200 Gln Glu Met

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1460)	sn Phe Trp Val Ser (1465	1470
Phe Leu Ala Gln 1475		ro Leu Ser Glu Glu 1 480	Phe Ser Tyr Tyr 1485
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1555	15		1565
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	1605	er Lys Asp Ser Ser I 1610	1615
1620)	er Ser Ile Arg Gln I 1625	1630
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Leu Ser Leu Pro 1650	Val Gly Cys Al 1655	la Val Glu Gly Ala 1 1660	Ile Met Asn Cys
Asn Ile Leu Met 1665	Tyr Asn Lys Le 1670	eu Ala Leu Ala Tyr N 1675	Met Pro Ser Ile 1680
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Ala Gly Gln Val 1700	Ile Cys Gly Va	al Pro Thr Arg Thr S 1705	
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374

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PCT/US01/23121 WO 02/08267

375

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Phe Ser Gln Ser Trp Glu Leu Gly Lys Phe Asn Glu Ser Arg Lys Leu

Ser Asn Gly Thr Leu Leu Glu Ile Ala Lys Ile Tyr Pro Met Asp Ala

Ser Phe Asp Ser Pro Glu Asp Val Pro Glu Asp Ile Ala Glu Asn Lys

Arg Tyr Lys Gly Ile Thr Gly Phe Thr Ile Ser Glu Val Ala Glu Gln

Val Lys Lys Asp Phe Gly His Ile Asp Ile Leu Val His Ser Leu Ala

Asn Ser Pro Glu Ile Ser Lys Ser Leu Leu Glu Thr Ser Arg Lys Gly

Tyr Leu Ala Ala Leu Ser Ala Ser Ser Tyr Ser Phe Val Ser Leu Leu 150 155

Ser His Phe Gly Ser Ile Met Asn Arg Gly Gly Ser Thr Ile Ser Leu

Thr Tyr Leu Ala Ser Met Arg Ala Val Pro Gly Tyr Gly Gly Met

Ser Ser Ala Lys Ala Ala Leu Glu Ser Asp Thr Lys Thr Leu Ala Trp

Glu Ala Gly Arg Arg Trp Gly Ile Arg Val Asn Thr Ile Ser Ala Gly

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Glu Gln Val Gly Ala Val Ala Ala Phe Leu Ala Ser Pro Leu Ala Ser

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376

295

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Ile Pro Tyr Tyr Thr Val Ser Phe Ser Lys Glu Tyr Lys Glu Arg Val 65 70 75 80

Phe Ser Arg Phe Leu Arg Glu Tyr Ala Asn Gly Tyr Thr Pro Asn Pro 85 90

Asp Val Leu Cys Asn Arg Glu Ile Lys Phe Asp Leu Leu Gln Lys Lys 100 105 110

Val Arg Glu Leu Lys Gly Asp Phe Leu Ala Thr Gly His Tyr Cys Arg 115 120 125

Gly Gly Ala Asp Gly Thr Gly Leu Ser Arg Gly Ile Asp Pro Asn Lys 130 135 140

Asp Gln Ser Tyr Phe Leu Cys Gly Thr Pro Lys Asp Ala Leu Ser Asn 145 150 155 160

Val Leu Phe Pro Leu Gly Gly Met Tyr Lys Thr Glu Val Arg Arg Ile 165 170 175

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Gly Leu Asn Ile Gly Gly Met Glu Lys Pro Cys Tyr Val Leu Ser Lys 245 250 255

Asn Met Glu Lys Asn Ile Val Tyr Ile Val Arg Gly Glu Asp His Pro 260 265 270

Leu Leu Tyr Arg Gln Glu Leu Leu Ala Lys Glu Leu Asn Trp Phe Val 275 280 285

Pro Leu Gln Glu Pro Met Ile Cys Ser Ala Lys Val Arg Tyr Arg Ser 290 295 300 379

Pro Asp Glu Lys Cys Ser Val Tyr Pro Leu Glu Asp Gly Thr Val Lys 305 310 315 320

Ala Phe Tyr Gln Gly Asp Ile Cys Leu Gly Gly Gly Val Ile Glu Val 340 $$ 345 $$ 350

Pro Met Ile His Gln Leu 355